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(54) Title: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

(57) Abstract:

NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

1. TECHNICAL FIELD

The present invention provides novel polynucleotides and proteins encoded by such polynucleotides, along with uses for these polynucleotides and proteins, for example in therapeutic, diagnostic and research methods.

2. BACKGROUND

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Technology aimed at the discovery of protein factors (including *e.g.*, cytokines, such as lymphokines, interferons, CSFs, chemokines, and interleukins) has matured rapidly over the past decade. The now routine hybridization cloning and expression cloning techniques clone novel polynucleotides "directly" in the sense that they rely on information directly related to the discovered protein (*i.e.*, partial DNA/amino acid sequence of the protein in the case of hybridization cloning; activity of the protein in the case of expression cloning). More recent "indirect" cloning techniques such as signal sequence cloning, which isolates DNA sequences based on the presence of a now well-recognized secretory leader sequence motif, as well as various PCR-based or low stringency hybridization-based cloning techniques, have advanced the state of the art by making available large numbers of DNA/amino acid sequences for proteins that are known to have biological activity, for example, by virtue of their secreted nature in the case of leader sequence cloning, by virtue of their cell or tissue source in the case of PCR-based techniques, or by virtue of structural similarity to other genes of known biological activity.

Identified polynucleotide and polypeptide sequences have numerous applications in, for example, diagnostics, forensics, gene mapping; identification of mutations responsible for genetic disorders or other traits, to assess biodiversity, and to produce many other types of data and products dependent on DNA and amino acid sequences.

3. SUMMARY OF THE INVENTION

The compositions of the present invention include novel isolated polypeptides, novel isolated polynucleotides encoding such polypeptides, including recombinant DNA molecules, cloned genes or degenerate variants thereof, especially naturally occurring variants such as allelic variants, antisense polynucleotide molecules, and antibodies that specifically recognize one or more epitopes present on such polypeptides, as well as hybridomas producing such antibodies.

The compositions of the present invention additionally include vectors, including expression vectors, containing the polynucleotides of the invention, cells genetically engineered to contain such polynucleotides and cells genetically engineered to express such polynucleotides.

The present invention relates to a collection or library of at least one novel nucleic acid sequence assembled from expressed sequence tags (ESTs) isolated mainly by sequencing by hybridization (SBH), and in some cases, sequences obtained from one or more public databases. The invention relates also to the proteins encoded by such polynucleotides, along with therapeutic, diagnostic and research utilities for these polynucleotides and proteins. These nucleic acid sequences are designated as SEQ ID NO: 1-1009. The polypeptides sequences are designated SEQ ID NO: 1010-2018. The nucleic acids and polypeptides are provided in the Sequence Listing. In the nucleic acids provided in the Sequence Listing, A is adenosine; C is cytosine; G is guanine; T is thymine; and N is any of the four bases. In the amino acids provided in the Sequence Listing, * corresponds to the stop codon.

The nucleic acid sequences of the present invention also include, nucleic acid sequences that hybridize to the complement of SEQ ID NO:1-1009 under stringent hybridization conditions; nucleic acid sequences which are allelic variants or species homologues of any of the nucleic acid sequences recited above, or nucleic acid sequences that encode a peptide comprising a specific domain or truncation of the peptides encoded by SEQ ID NO:1-1009. A polynucleotide comprising a nucleotide sequence having at least 90% identity to an identifying sequence of SEQ ID NO:1-1009 or a degenerate variant or fragment thereof. The identifying sequence can be 100 base pairs in length.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO:1-1009 . The sequence information can be a segment of any one of SEQ ID NO:1-1009 that uniquely identifies or represents the sequence information of SEQ ID NO:1-1009.

A collection as used in this application can be a collection of only one polynucleotide. The collection of sequence information or identifying information of each sequence can be provided on a nucleic acid array. In one embodiment, segments of sequence information is provided on a nucleic acid array to detect the polynucleotide that contains the segment. The array can be designed to detect full-match or mismatch to the polynucleotide that contains the segment. The collection can also be provided in a computer-readable format.

This invention also includes the reverse or direct complement of any of the nucleic acid sequences recited above; cloning or expression vectors containing the nucleic acid sequences; and host cells or organisms transformed with these expression vectors. Nucleic acid sequences (or their reverse or direct complements) according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology, such as use as hybridization probes, use as primers for PCR, use in an array, use in computer-readablemedia, use in sequencing

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full-length genes, use for chromosome and gene mapping, use in the recombinant production of protein, and use in the generation of anti-sense DNA or RNA, their chemical analogs and the like.

In a preferred embodiment, the nucleic acid sequences of SEQ ID NO:1-1009 or novel segments or parts of the nucleic acids of the invention are used as primers in expression assays that are well known in the art. In a particularly preferred embodiment, the nucleic acid sequences of SEQ ID NO:1-1009 or novel segments or parts of the nucleic acids provided herein are used in diagnostics for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The isolated polynucleotides of the invention include, but are not limited to, a polynucleotide comprising any one of the nucleotide sequences set forth in SEQ ID NO:1-1009; a polynucleotide comprising any of the full length protein coding sequences of SEQ ID NO:1 - 1009; and a polynucleotide comprising any of the nucleotide sequences of the mature protein coding sequences of SEQ ID NO: 1- 1009. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent hybridization conditions to (a) the complement of any one of the nucleotide sequences set forth in SEQ ID NO:1-1009; (b) a nucleotide sequence encoding any one of the amino acid sequences set forth in the Sequence Listing (e.g., SEQ ID NO: 1010-2018); (c) a polynucleotide which is an allelic variant of any polynucleotides recited above; (d) a polynucleotide which encodes a species homolog (e.g. orthologs) of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of any of the polypeptides comprising an amino acid sequence set forth in the Sequence Listing.

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising any of the amino acid sequences set forth in the Sequence Listing; or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides with biological activity that are encoded by (a) any of the polynucleotides having a nucleotide sequence set forth in SEQ ID NO:1-1009; or (b) polynucleotides that hybridize to the complement of the polynucleotides of (a) under stringent hybridization conditions. Biologically or immunologically active variants of any of the polypeptide sequences in the Sequence Listing, and "substantial equivalents" thereof (e.g., with at least about 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or 99% amino acid sequence identity) that preferably retain biological activity are also contemplated. The polypeptides of the invention may be wholly or partially chemically synthesized but are preferably produced by recombinant means using the genetically engineered cells (e.g. host cells) of the invention.

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The invention also provides compositions comprising a polypeptide of the invention. Polypeptide compositions of the invention may further comprise an acceptable carrier, such as a hydrophilic, *e.g.*, pharmaceutically acceptable, carrier.

The invention also provides host cells transformed or transfected with a polynucleotide of the invention.

The invention also relates to methods for producing a polypeptide of the invention comprising growing a culture of the host cells of the invention in a suitable culture medium under conditions permitting expression of the desired polypeptide, and purifying the polypeptide from the culture or from the host cells. Preferred embodiments include those in which the protein produced by such process is a mature form of the protein.

Polynucleotides according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology. These techniques include use as hybridization probes, use as oligomers, or primers, for PCR, use for chromosome and gene mapping, use in the recombinant production of protein, and use in generation of anti-sense DNA or RNA, their chemical analogs and the like. For example, when the expression of an mRNA is largely restricted to a particular cell or tissue type, polynucleotides of the invention can be used as hybridization probes to detect the presence of the particular cell or tissue mRNA in a sample using, *e.g.*, *in situ* hybridization.

In other exemplary embodiments, the polynucleotides are used in diagnostics as expressed sequence tags for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The polypeptides according to the invention can be used in a variety of conventional procedures and methods that are currently applied to other proteins. For example, a polypeptide of the invention can be used to generate an antibody that specifically binds the polypeptide. Such antibodies, particularly monoclonal antibodies, are useful for detecting or quantitating the polypeptide in tissue. The polypeptides of the invention can also be used as molecular weight markers, and as a food supplement.

Methods are also provided for preventing, treating, or ameliorating a medical condition which comprises the step of administering to a mammalian subject a therapeutically effective amount of a composition comprising a polypeptide of the present invention and a pharmaceutically acceptable carrier.

In particular, the polypeptides and polynucleotides of the invention can be utilized, for example, in methods for the prevention and/or treatment of disorders involving aberrant protein expression or biological activity.

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The present invention further relates to methods for detecting the presence of the polynucleotides or polypeptides of the invention in a sample. Such methods can, for example, be utilized as part of prognostic and diagnostic evaluation of disorders as recited herein and for the identification of subjects exhibiting a predisposition to such conditions. The invention provides a method for detecting the polynucleotides of the invention in a sample, comprising contacting the sample with a compound that binds to and forms a complex with the polynucleotide of interest for a period sufficient to form the complex and under conditions sufficient to form a complex and detecting the complex such that if a complex is detected, the polynucleotide of interest is detected. The invention also provides a method for detecting the polypeptides of the invention in a sample comprising contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex and detecting the formation of the complex such that if a complex is formed, the polypeptide is detected.

The invention also provides kits comprising polynucleotide probes and/or monoclonal antibodies, and optionally quantitative standards, for carrying out methods of the invention. Furthermore, the invention provides methods for evaluating the efficacy of drugs, and monitoring the progress of patients, involved in clinical trials for the treatment of disorders as recited above.

The invention also provides methods for the identification of compounds that modulate (*i.e.*, increase or decrease) the expression or activity of the polynucleotides and/or polypeptides of the invention. Such methods can be utilized, for example, for the identification of compounds that can ameliorate symptoms of disorders as recited herein. Such methods can include, but are not limited to, assays for identifying compounds and other substances that interact with (*e.g.*, bind to) the polypeptides of the invention. The invention provides a method for identifying a compound that binds to the polypeptides of the invention comprising contacting the compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and detecting the complex by detecting the reporter gene sequence expression such that if expression of the reporter gene is detected the compound the binds to a polypeptide of the invention is identified.

The methods of the invention also provides methods for treatment which involve the administration of the polynucleotides or polypeptides of the invention to individuals exhibiting symptoms or tendencies. In addition, the invention encompasses methods for treating diseases or disorders as recited herein comprising administering compounds and other substances that modulate the overall activity of the target gene products. Compounds and other substances can

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effect such modulation either on the level of target gene/protein expression or target protein activity.

The polypeptides of the present invention and the polynucleotides encoding them are also useful for the same functions known to one of skill in the art as the polypeptides and polynucleotides to which they have homology (set forth in Table 2). If no homology is set forth for a sequence, then the polypeptides and polynucleotides of the present invention are useful for a variety of applications, as described herein, including use in arrays for detection.

4. DETAILED DESCRIPTION OF THE INVENTION

4.1 DEFINITIONS

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It must be noted that as used herein and in the appended claims, the singular forms "a", "an" and "the" include plural references unless the context clearly dictates otherwise.

The term "active" refers to those forms of the polypeptide which retain the biologic and/or immunologic activities of any naturally occurring polypeptide. According to the invention, the terms "biologically active" or "biological activity" refer to a protein or peptide having structural, regulatory or biochemical functions of a naturally occurring molecule. Likewise "immunologically active" or "immunological activity" refers to the capability of the natural, recombinant or synthetic polypeptide to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies.

The term "activated cells" as used in this application are those cells which are engaged in extracellular or intracellular membrane trafficking, including the export of secretory or enzymatic molecules as part of a normal or disease process.

The terms "complementary" or "complementarity" refer to the natural binding of polynucleotides by base pairing. For example, the sequence 5'-AGT-3' binds to the complementary sequence 3'-TCA-5'. Complementarity between two single-stranded molecules may be "partial" such that only some of the nucleic acids bind or it may be "complete" such that total complementarity exists between the single stranded molecules. The degree of complementarity between the nucleic acid strands has significant effects on the efficiency and strength of the hybridization between the nucleic acid strands.

The term "embryonic stem cells (ES)" refers to a cell that can give rise to many differentiated cell types in an embryo or an adult, including the germ cells. The term "germ line stem cells (GSCs)" refers to stem cells derived from primordial stem cells that provide a steady and continuous source of germ cells for the production of gametes. The term "primordial germ

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cells (PGCs)" refers to a small population of cells set aside from other cell lineages particularly from the yolk sac, mesenteries, or gonadal ridges during embryogenesis that have the potential to differentiate into germ cells and other cells. PGCs are the source from which GSCs and ES cells are derived The PGCs, the GSCs and the ES cells are capable of self-renewal. Thus these cells not only populate the germ line and give rise to a plurality of terminally differentiated cells that comprise the adult specialized organs, but are able to regenerate themselves.

The term "expression modulating fragment," EMF, means a series of nucleotides which modulates the expression of an operably linked ORF or another EMF.

As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are nucleic acid fragments which induce the expression of an operably linked ORF in response to a specific regulatory factor or physiological event.

The terms "nucleotide sequence" or "nucleic acid" or "polynucleotide" or "oligonculeotide" are used interchangeably and refer to a heteropolymer of nucleotides or the sequence of these nucleotides. These phrases also refer to DNA or RNA of genomic or synthetic origin which may be single-stranded or double-stranded and may represent the sense or the antisense strand, to peptide nucleic acid (PNA) or to any DNA-like or RNA-like material. In the sequences herein A is adenine, C is cytosine, T is thymine, G is guanine and N is A, C, G or T (U). It is contemplated that where the polynucleotide is RNA, the T (thymine) in the sequences provided herein is substituted with U (uracil). Generally, nucleic acid segments provided by this invention may be assembled from fragments of the genome and short oligonucleotide linkers, or from a series of oligonucleotides, or from individual nucleotides, to provide a synthetic nucleic acid which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon, or a eukaryotic gene.

The terms "oligonucleotide fragment" or a "polynucleotide fragment", "portion," or "segment" or "probe" or "primer" are used interchangeably and refer to a sequence of nucleotide residues which are at least about 5 nucleotides, more preferably at least about 7 nucleotides, more preferably at least about 9 nucleotides, more preferably at least about 11 nucleotides and most preferably at least about 17 nucleotides. The fragment is preferably less than about 500 nucleotides, preferably less than about 200 nucleotides, more preferably less than about 100 nucleotides, more preferably less than about 50 nucleotides and most preferably less than 30 nucleotides. Preferably the probe is from about 6 nucleotides to about 200 nucleotides. preferably from about 15 to about 50 nucleotides, more preferably from about 17 to 30 nucleotides and most preferably from about 20 to 25 nucleotides. Preferably the fragments can

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be used in polymerase chain reaction (PCR), various hybridization procedures or microarray procedures to identify or amplify identical or related parts of mRNA or DNA molecules. A fragment or segment may uniquely identify each polynucleotide sequence of the present invention. Preferably the fragment comprises a sequence substantially similar to any one of SEQ ID NOs:1-1009.

Probes may, for example, be used to determine whether specific mRNA molecules are present in a cell or tissue or to isolate similar nucleic acid sequences from chromosomal DNA as described by Walsh et al. (Walsh, P.S. et al., 1992, PCR Methods Appl 1:241-250). They may be labeled by nick translation, Klenow fill-in reaction, PCR, or other methods well known in the art. Probes of the present invention, their preparation and/or labeling are elaborated in Sambrook, J. et al., 1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY; or Ausubel, F.M. et al., 1989, Current Protocols in Molecular Biology, John Wiley & Sons, New York NY, both of which are incorporated herein by reference in their entirety.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO:1-1009. The sequence information can be a segment of any one of SEQ ID NO:1-1009 that uniquely identifies or represents the sequence information of that sequence of SEQ ID NO:1-1009. One such segment can be a twenty-mer nucleic acid sequence because the probability that a twenty-mer is fully matched in the human genome is 1 in 300. In the human genome, there are three billion base pairs in one set of chromosomes. Because 4²⁰ possible twenty-mers exist, there are 300 times more twenty-mers than there are base pairs in a set of human chromosomes. Using the same analysis, the probability for a seventeen-mer to be fully matched in the human genome is approximately 1 in 5. When these segments are used in arrays for expression studies, fifteen-mer segments can be used. The probability that the fifteen-mer is fully matched in the expressed sequences is also approximately one in five because expressed sequences comprise less than approximately 5% of the entire genome sequence.

Similarly, when using sequence information for detecting a single mismatch, a segment can be a twenty-five mer. The probability that the twenty-five mer would appear in a human genome with a single mismatch is calculated by multiplying the probability for a full match $(1 \div 4^{25})$ times the increased probability for mismatch at each nucleotide position (3×25) . The probability that an eighteen mer with a single mismatch can be detected in an array for expression studies is approximately one in five. The probability that a twenty-mer with a single mismatch can be detected in a human genome is approximately one in five.

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The term "open reading frame," ORF, means a series of nucleotide triplets coding for amino acids without any termination codons and is a sequence translatable into protein.

The terms "operably linked" or "operably associated" refer to functionally related nucleic acid sequences. For example, a promoter is operably associated or operably linked with a coding sequence if the promoter controls the transcription of the coding sequence. While operably linked nucleic acid sequences can be contiguous and in the same reading frame, certain genetic elements *e.g.* repressor genes are not contiguously linked to the coding sequence but still control transcription/translation of the coding sequence.

The term "pluripotent" refers to the capability of a cell to differentiate into a number of differentiated cell types that are present in an adult organism. A pluripotent cell is restricted in its differentiation capability in comparison to a totipotent cell.

The terms "polypeptide" or "peptide" or "amino acid sequence" refer to an oligopeptide, peptide, polypeptide or protein sequence or fragment thereof and to naturally occurring or synthetic molecules. A polypeptide "fragment," "portion," or "segment" is a stretch of amino acid residues of at least about 5 amino acids, preferably at least about 7 amino acids, more preferably at least about 9 amino acids and most preferably at least about 17 or more amino acids. The peptide preferably is not greater than about 200 amino acids, more preferably less than 150 amino acids and most preferably less than 100 amino acids. Preferably the peptide is from about 5 to about 200 amino acids. To be active, any polypeptide must have sufficient length to display biological and/or immunological activity.

The term "naturally occurring polypeptide" refers to polypeptides produced by cells that have not been genetically engineered and specifically contemplates various polypeptides arising from post-translational modifications of the polypeptide including, but not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation and acylation.

The term "translated protein coding portion" means a sequence which encodes for the full length protein which may include any leader sequence or any processing sequence.

The term "mature protein coding sequence" means a sequence which encodes a peptide or protein without a signal or leader sequence. The "mature protein portion" means that portion of the protein which does not include a signal or leader sequence. The peptide may have been produced by processing in the cell which removes any leader/signal sequence. The mature protein portion may or may not include the initial methionine residue. The methionine residue may be removed from the protein during processing in the cell. The peptide may be produced synthetically or the protein may have been produced using a polynucleotide only encoding for the mature protein coding sequence.

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The term "derivative" refers to polypeptides chemically modified by such techniques as ubiquitination, labeling (e.g., with radionuclides or various enzymes), covalent polymer attachment such as pegylation (derivatization with polyethylene glycol) and insertion or substitution by chemical synthesis of amino acids such as ornithine, which do not normally occur in human proteins.

The term "variant" (or "analog") refers to any polypeptide differing from naturally occurring polypeptides by amino acid insertions, deletions, and substitutions, created using, e.g., recombinant DNA techniques. Guidance in determining which amino acid residues may be replaced, added or deleted without abolishing activities of interest, may be found by comparing the sequence of the particular polypeptide with that of homologous peptides and minimizing the number of amino acid sequence changes made in regions of high homology (conserved regions) or by replacing amino acids with consensus sequence.

Alternatively, recombinant variants encoding these same or similar polypeptides may be synthesized or selected by making use of the "redundancy" in the genetic code. Various codon substitutions, such as the silent changes which produce various restriction sites, may be introduced to optimize cloning into a plasmid or viral vector or expression in a particular prokaryotic or eukaryotic system. Mutations in the polynucleotide sequence may be reflected in the polypeptide or domains of other peptides added to the polypeptide to modify the properties of any part of the polypeptide, to change characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate.

Preferably, amino acid "substitutions" are the result of replacing one amino acid with another amino acid having similar structural and/or chemical properties, *i.e.*, conservative amino acid replacements. "Conservative" amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues involved. For example, nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan, and methionine; polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine; positively charged (basic) amino acids include arginine, lysine, and histidine; and negatively charged (acidic) amino acids include aspartic acid and glutamic acid. "Insertions" or "deletions" are preferably in the range of about 1 to 20 amino acids, more preferably 1 to 10 amino acids. The variation allowed may be experimentally determined by systematically making insertions, deletions, or substitutions of amino acids in a polypeptide molecule using recombinant DNA techniques and assaying the resulting recombinant variants for activity.

Alternatively, where alteration of function is desired, insertions, deletions or non-conservative alterations can be engineered to produce altered polypeptides. Such alterations

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can, for example, alter one or more of the biological functions or biochemical characteristics of the polypeptides of the invention. For example, such alterations may change polypeptide characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate. Further, such alterations can be selected so as to generate polypeptides that are better suited for expression, scale up and the like in the host cells chosen for expression. For example, cysteine residues can be deleted or substituted with another amino acid residue in order to eliminate disulfide bridges.

The terms "purified" or "substantially purified" as used herein denotes that the indicated nucleic acid or polypeptide is present in the substantial absence of other biological macromolecules, e.g., polynucleotides, proteins, and the like. In one embodiment, the polynucleotide or polypeptide is purified such that it constitutes at least 95% by weight, more preferably at least 99% by weight, of the indicated biological macromolecules present (but water, buffers, and other small molecules, especially molecules having a molecular weight of less than 1000 daltons, can be present).

The term "isolated" as used herein refers to a nucleic acid or polypeptide separated from at least one other component (e.g., nucleic acid or polypeptide) present with the nucleic acid or polypeptide in its natural source. In one embodiment, the nucleic acid or polypeptide is found in the presence of (if anything) only a solvent, buffer, ion, or other component normally present in a solution of the same. The terms "isolated" and "purified" do not encompass nucleic acids or polypeptides present in their natural source.

The term "recombinant," when used herein to refer to a polypeptide or protein, means that a polypeptide or protein is derived from recombinant (e.g., microbial, insect, or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (e.g., yeast) expression systems. As a product, "recombinant microbial" defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, e.g., E. coli, will be free of glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern in general different from those expressed in mammalian cells.

The term "recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. An expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers, (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3)

35 appropriate transcription initiation and termination sequences. Structural units intended for use

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in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an amino terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

The term "recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extrachromosomally. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed. This term also means host cells which have stably integrated a recombinant genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers. Recombinant expression systems as defined herein will express polypeptides or proteins endogenous to the cell upon induction of the regulatory elements linked to the endogenous DNA segment or gene to be expressed. The cells can be prokaryotic or eukaryotic.

The term "secreted" includes a protein that is transported across or through a membrane, including transport as a result of signal sequences in its amino acid sequence when it is expressed in a suitable host cell. "Secreted" proteins include without limitation proteins secreted wholly (e.g., soluble proteins) or partially (e.g., receptors) from the cell in which they are expressed. "Secreted" proteins also include without limitation proteins that are transported across the membrane of the endoplasmic reticulum. "Secreted" proteins are also intended to include proteins containing non-typical signal sequences (e.g. Interleukin-1 Beta, see Krasney, P.A. and Young, P.R. (1992) Cytokine 4(2):134-143) and factors released from damaged cells (e.g. Interleukin-1 Receptor Antagonist, see Arend, W.P. et. al. (1998) Annu. Rev. Immunol. 16:27-55)

Where desired, an expression vector may be designed to contain a "signal or leader sequence" which will direct the polypeptide through the membrane of a cell. Such a sequence may be naturally present on the polypeptides of the present invention or provided from heterologous protein sources by recombinant DNA techniques.

The term "stringent" is used to refer to conditions that are commonly understood in the art as stringent. Stringent conditions can include highly stringent conditions (*i.e.*, hybridization to filter-bound DNA in 0.5 M NaHPO₄, 7% sodium dodecyl sulfate (SDS), 1 mM EDTA at 65°C, and washing in 0.1X SSC/0.1% SDS at 68°C), and moderately stringent conditions (*i.e.*, washing in 0.2X SSC/0.1% SDS at 42°C). Other exemplary hybridization conditions are described herein in the examples.

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In instances of hybridization of deoxyoligonucleotides, additional exemplary stringent hybridization conditions include washing in 6X SSC/0.05% sodium pyrophosphate at 37°C (for 14-base oligonucleotides), 48°C (for 17-base oligos), 55°C (for 20-base oligonucleotides), and 60°C (for 23-base oligonucleotides).

As used herein, "substantially equivalent" can refer both to nucleotide and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between the reference and subject sequences. Typically, such a substantially equivalent sequence varies from one of those listed herein by no more than about 35% (i.e., the number of individual residue substitutions, additions, and/or deletions in a substantially equivalent sequence, as compared to the corresponding reference sequence, divided by the total number of residues in the substantially equivalent sequence is about 0.35 or less). Such a sequence is said to have 65% sequence identity to the listed sequence. In one embodiment, a substantially equivalent, e.g., mutant, sequence of the invention varies from a listed sequence by no more than 30% (70% sequence identity); in a variation of this embodiment, by no more than 25% (75% sequence identity); and in a further variation of this embodiment, by no more than 20% (80% sequence identity) and in a further variation of this embodiment, by no more than 10% (90% sequence identity) and in a further variation of this embodiment, by no more that 5% (95% sequence identity). Substantially equivalent, e.g., mutant, amino acid sequences according to the invention preferably have at least 80% sequence identity with a listed amino acid sequence, more preferably at least 85% sequence identity, more preferably at least 90% sequence identity, more preferably at least 95% identity, more preferably at least 98% identity, and most preferably at least 99% identity. Substantially equivalent nucleotide sequences of the invention can have lower percent sequence identities, taking into account, for example, the redundancy or degeneracy of the genetic code. Preferably, nucleotide sequence has at least about 65% identity, more preferably at least about 75% identity, more preferably at least about 80% sequence identity, more preferably at least about 85% sequence identity, more preferably at least about 90% sequence identity, and most preferably at least about 95% identity, more preferably at least about 98% sequence identity, and most preferably at least about 99% sequence identity. For the purposes of the present invention, sequences having substantially equivalent biological activity and substantially equivalent expression characteristics are considered substantially equivalent. For the purposes of determining equivalence, truncation of the mature sequence (e.g., via a mutation which creates a spurious stop codon) should be disregarded. Sequence identity may be determined, e.g., using the Jotun Hein method (Hein, J.

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(1990) Methods Enzymol. 183:626-645). Identity between sequences can also be determined by other methods known in the art, e.g. by varying hybridization conditions.

The term "totipotent" refers to the capability of a cell to differentiate into all of the cell types of an adult organism.

The term "transformation" means introducing DNA into a suitable host cell so that the DNA is replicable, either as an extrachromosomal element, or by chromosomal integration. The term "transfection" refers to the taking up of an expression vector by a suitable host cell, whether or not any coding sequences are in fact expressed. The term "infection" refers to the introduction of nucleic acids into a suitable host cell by use of a virus or viral vector.

As used herein, an "uptake modulating fragment," UMF, means a series of nucleotides which mediate the uptake of a linked DNA fragment into a cell. UMFs can be readily identified using known UMFs as a target sequence or target motif with the computer-based systems described below. The presence and activity of a UMF can be confirmed by attaching the suspected UMF to a marker sequence. The resulting nucleic acid molecule is then incubated with an appropriate host under appropriate conditions and the uptake of the marker sequence is determined. As described above, a UMF will increase the frequency of uptake of a linked marker sequence.

Each of the above terms is meant to encompass all that is described for each, unless the context dictates otherwise.

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4.2 NUCLEIC ACIDS OF THE INVENTION

Nucleotide sequences of the invention are set forth in the Sequence Listing.

The isolated polynucleotides of the invention include a polynucleotide comprising the nucleotide sequences of SEQ ID NO:1-1009; a polynucleotide encoding any one of the peptide sequences of SEQ ID NO:1010-2018; and a polynucleotide comprising the nucleotide sequence encoding the mature protein coding sequence of the polypeptides of any one of SEQ ID NO:1010-2018. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent conditions to (a) the complement of any of the nucleotides sequences of SEQ ID NO:1-1009; (b) nucleotide sequences encoding any one of the amino acid sequences set forth in the Sequence Listing; (c) a polynucleotide which is an allelic variant of any polynucleotide recited above; (d) a polynucleotide which encodes a species homolog of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of the polypeptides of SEQ ID NO: 1010-2018.

Domains of interest may depend on the nature of the encoded polypeptide; e.g., domains in receptor-like polypeptides include ligand-binding, extracellular, transmembrane, or cytoplasmic

domains, or combinations thereof; domains in immunoglobulin-like proteins include the variable . immunoglobulin-like domains; domains in enzyme-like polypeptides include catalytic and substrate binding domains; and domains in ligand polypeptides include receptor-binding domains.

The polynucleotides of the invention include naturally occurring or wholly or partially synthetic DNA, e.g., cDNA and genomic DNA, and RNA, e.g., mRNA. The polynucleotides may include all of the coding region of the cDNA or may represent a portion of the coding region of the cDNA.

The present invention also provides genes corresponding to the cDNA sequences disclosed herein. The corresponding genes can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include the preparation of probes or primers from the disclosed sequence information for identification and/or amplification of genes in appropriate genomic libraries or other sources of genomic materials. Further 5' and 3' sequence can be obtained using methods known in the art. For example, full length cDNA or genomic DNA that corresponds to any of the polynucleotides of SEQ ID NO:1-1009 can be obtained by screening appropriate cDNA or genomic DNA libraries under suitable hybridization conditions using any of the polynucleotides of SEQ ID NO:1-1009 or a portion thereof as a probe. Alternatively, the polynucleotides of SEQ ID NO:1-1009 may be used as the basis for suitable primer(s) that allow identification and/or amplification of genes in appropriate genomic DNA or cDNA libraries.

The nucleic acid sequences of the invention can be assembled from ESTs and sequences (including cDNA and genomic sequences) obtained from one or more public databases, such as dbEST, gbpri, and UniGene. The EST sequences can provide identifying sequence information, representative fragment or segment information, or novel segment information for the full-length gene.

The polynucleotides of the invention also provide polynucleotides including nucleotide sequences that are substantially equivalent to the polynucleotides recited above. Polynucleotides according to the invention can have, *e.g.*, at least about 65%, at least about 70%, at least about 75%, at least about 80%, 81%, 82%, 83%, 84%, more typically at least about 85%, 86%, 87%, 88%, 89%, more typically at least about 90%, 91%, 92%, 93%, 94%, and even more typically at least about 95%, 96%, 97%, 98%, 99%, sequence identity to a polynucleotide recited above.

Included within the scope of the nucleic acid sequences of the invention are nucleic acid sequence fragments that hybridize under stringent conditions to any of the nucleotide sequences of SEQ ID NO:1-1009, or complements thereof, which fragment is greater than about 5 nucleotides, preferably 7 nucleotides, more preferably greater than 9 nucleotides and most preferably greater than 17 nucleotides. Fragments of, e.g. 15, 17, or 20 nucleotides or more that

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are selective for (i.e. specifically hybridize to any one of the polynucleotides of the invention) are contemplated. Probes capable of specifically hybridizing to a polynucleotide can differentiate polynucleotide sequences of the invention from other polynucleotide sequences in the same family of genes or can differentiate human genes from genes of other species, and are preferably based on unique nucleotide sequences.

The sequences falling within the scope of the present invention are not limited to these specific sequences, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequence provided SEQ ID NO:1-1009, a representative fragment thereof, or a nucleotide sequence at least 90% identical, preferably 95% identical, to SEQ ID NO:1-1009 with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another codon that encodes the same amino acid is expressly contemplated.

The nearest neighbor or homology result for the nucleic acids of the present invention, including SEQ ID NO:1-1009, can be obtained by searching a database using an algorithm or a program. Preferably, a BLAST which stands for Basic Local Alignment Search Tool is used to search for local sequence alignments (Altshul, S.F. J Mol. Evol. 36 290-300 (1993) and Altschul S.F. et al. J. Mol. Biol. 21:403-410 (1990)). Alternatively a FASTA version 3 search against Genpept, using Fastxy algorithm.

Species homologs (or orthologs) of the disclosed polynucleotides and proteins are also provided by the present invention. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source from the desired species.

The invention also encompasses allelic variants of the disclosed polynucleotides or proteins; that is, naturally-occurring alternative forms of the isolated polynucleotide which also encode proteins which are identical, homologous or related to that encoded by the polynucleotides.

The nucleic acid sequences of the invention are further directed to sequences which encode variants of the described nucleic acids. These amino acid sequence variants may be prepared by methods known in the art by introducing appropriate nucleotide changes into a native or variant polynucleotide. There are two variables in the construction of amino acid sequence variants: the location of the mutation and the nature of the mutation. Nucleic acids encoding the amino acid sequence variants are preferably constructed by mutating the polynucleotide to encode an amino acid sequence that does not occur in nature. These nucleic

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acid alterations can be made at sites that differ in the nucleic acids from different species (variable positions) or in highly conserved regions (constant regions). Sites at such locations will typically be modified in series, *e.g.*, by substituting first with conservative choices (*e.g.*, hydrophobic amino acid to a different hydrophobic amino acid) and then with more distant choices (*e.g.*, hydrophobic amino acid to a charged amino acid), and then deletions or insertions may be made at the target site. Amino acid sequence deletions generally range from about 1 to 30 residues, preferably about 1 to 10 residues, and are typically contiguous. Amino acid insertions include amino- and/or carboxyl-terminal fusions ranging in length from one to one hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. Intrasequence insertions may range generally from about 1 to 10 amino residues, preferably from 1 to 5 residues. Examples of terminal insertions include the heterologous signal sequences necessary for secretion or for intracellular targeting in different host cells and sequences such as FLAG or poly-histidine sequences useful for purifying the expressed protein.

In a preferred method, polynucleotides encoding the novel amino acid sequences are changed via site-directed mutagenesis. This method uses oligonucleotide sequences to alter a polynucleotide to encode the desired amino acid variant, as well as sufficient adjacent nucleotides on both sides of the changed amino acid to form a stable duplex on either side of the site of being changed. In general, the techniques of site-directed mutagenesis are well known to those of skill in the art and this technique is exemplified by publications such as, Edelman et al., *DNA* 2:183 (1983). A versatile and efficient method for producing site-specific changes in a polynucleotide sequence was published by Zoller and Smith, *Nucleic Acids Res.* 10:6487-6500 (1982). PCR may also be used to create amino acid sequence variants of the novel nucleic acids. When small amounts of template DNA are used as starting material, primer(s) that differs slightly in sequence from the corresponding region in the template DNA can generate the desired amino acid variant. PCR amplification results in a population of product DNA fragments that differ from the polynucleotide template encoding the polypeptide at the position specified by the primer. The product DNA fragments replace the corresponding region in the plasmid and this gives a polynucleotide encoding the desired amino acid variant.

A further technique for generating amino acid variants is the cassette mutagenesis technique described in Wells et al., *Gene* 34:315 (1985); and other mutagenesis techniques well known in the art, such as, for example, the techniques in Sambrook et al., supra, and *Current Protocols in Molecular Biology*, Ausubel et al. Due to the inherent degeneracy of the genetic code, other DNA sequences which encode substantially the same or a functionally equivalent amino acid sequence may be used in the practice of the invention for the cloning and expression

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of these novel nucleic acids. Such DNA sequences include those which are capable of hybridizing to the appropriate novel nucleic acid sequence under stringent conditions.

Polynucleotides encoding preferred polypeptide truncations of the invention can be used to generate polynucleotides encoding chimeric or fusion proteins comprising one or more domains of the invention and heterologous protein sequences.

The polynucleotides of the invention additionally include the complement of any of the polynucleotides recited above. The polynucleotide can be DNA (genomic, cDNA, amplified, or synthetic) or RNA. Methods and algorithms for obtaining such polynucleotides are well known to those of skill in the art and can include, for example, methods for determining hybridization conditions that can routinely isolate polynucleotides of the desired sequence identities.

In accordance with the invention, polynucleotide sequences comprising the mature protein coding sequences corresponding to any one of SEQ ID NO:1-1009, or functional equivalents thereof, may be used to generate recombinant DNA molecules that direct the expression of that nucleic acid, or a functional equivalent thereof, in appropriate host cells. Also included are the cDNA inserts of any of the clones identified herein.

A polynucleotide according to the invention can be joined to any of a variety of other nucleotide sequences by well-established recombinant DNA techniques (see Sambrook J et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY). Useful nucleotide sequences for joining to polynucleotides include an assortment of vectors, e.g., plasmids, cosmids, lambda phage derivatives, phagemids, and the like, that are well known in the art. Accordingly, the invention also provides a vector including a polynucleotide of the invention and a host cell containing the polynucleotide. In general, the vector contains an origin of replication functional in at least one organism, convenient restriction endonuclease sites, and a selectable marker for the host cell. Vectors according to the invention include expression vectors, replication vectors, probe generation vectors, and sequencing vectors. A host cell according to the invention can be a prokaryotic or eukaryotic cell and can be a unicellular organism or part of a multicellular organism.

The present invention further provides recombinant constructs comprising a nucleic acid having any of the nucleotide sequences of SEQ ID NO:1-1009 or a fragment thereof or any other polynucleotides of the invention. In one embodiment, the recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a nucleic acid having any of the nucleotide sequences of SEQ ID NO:1-1009 or a fragment thereof is inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF. Large numbers of suitable vectors and promoters are

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known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Bacterial: pBs, phagescript, PsiX174, pBluescript SK, pBs KS, pNH8a, pNH16a, pNH18a, pNH46a (Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (Pharmacia).

5 Eukaryotic: pWLneo, pSV2cat, pOG44, PXTI, pSG (Stratagene) pSVK3, pBPV, pMSG, pSVL (Pharmacia).

The isolated polynucleotide of the invention may be operably linked to an expression control sequence such as the pMT2 or pED expression vectors disclosed in Kaufman et al., *Nucleic Acids Res.* 19, 4485-4490 (1991), in order to produce the protein recombinantly. Many suitable expression control sequences are known in the art. General methods of expressing recombinant proteins are also known and are exemplified in R. Kaufman, *Methods in Enzymology* 185, 537-566 (1990). As defined herein "operably linked" means that the isolated polynucleotide of the invention and an expression control sequence are situated within a vector or cell in such a way that the protein is expressed by a host cell which has been transformed (transfected) with the ligated polynucleotide/expression control sequence.

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art. Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, e.g., the ampicillin resistance gene of E. coli and S. cerevisiae TRP1 gene, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), a-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an amino terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product. Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or

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more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and to, if desirable, provide amplification within the host. Suitable prokaryotic hosts for transformation include *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*, although others may also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (Promega Biotech, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed. Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is induced or derepressed by appropriate means (e.g., temperature shift or chemical induction) and cells are cultured for an additional period. Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

Polynucleotides of the invention can also be used to induce immune responses. For example, as described in Fan et al., *Nat. Biotech.* 17:870-872 (1999), incorporated herein by reference, nucleic acid sequences encoding a polypeptide may be used to generate antibodies against the encoded polypeptide following topical administration of naked plasmid DNA or following injection, and preferably intramuscular injection of the DNA. The nucleic acid sequences are preferably inserted in a recombinant expression vector and may be in the form of naked DNA.

25 **4.3 ANTISENSE**

Another aspect of the invention pertains to isolated antisense nucleic acid molecules that are hybridizable to or complementary to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NO:1-1009, or fragments, analogs or derivatives thereof. An "antisense" nucleic acid comprises a nucleotide sequence that is complementary to a "sense" nucleic acid encoding a protein, *e.g.*, complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence. In specific aspects, antisense nucleic acid molecules are provided that comprise a sequence complementary to at least about 10, 25, 50, 100, 250 or 500 nucleotides or an entire coding strand, or to only a portion thereof. Nucleic acid molecules encoding fragments, homologs, derivatives and analogs of a protein of any of SEQ ID

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NO:1010-2018 or antisense nucleic acids complementary to a nucleic acid sequence of SEQ ID NO:1-1009 are additionally provided.

In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence of the invention. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues. In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence of the invention. The term "noncoding region" refers to 5' and 3' sequences which flank the coding region that are not translated into amino acids (*i.e.*, also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding a nucleic acid disclosed herein (e.g., SEQ ID NO:1-1009), antisense nucleic acids of the invention can be designed according to the rules of Watson and Crick or Hoogsteen base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of a mRNA, but more preferably is an oligonucleotide that is antisense to only a portion of the coding or noncoding region of a mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of a mRNA. An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis or enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used.

Examples of modified nucleotides that can be used to generate the antisense nucleic acid
include: 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine,
4-acetylcytosine, 5-(carboxyhydroxylmethyl) uracil, 5-carboxymethylaminomethyl2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine,
inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine,
2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine,
7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil,
beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil,
2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil,
queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil,
uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil,
35 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the

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antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (*i.e.*, RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated in situ such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding a protein according to the invention to thereby inhibit expression of the protein, e.g., by inhibiting transcription and/or translation. The hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule that binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site. Alternatively, antisense nucleic acid molecules can be modified to target selected cells and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface, e.g., by linking the antisense nucleic acid molecules to peptides or antibodies that bind to cell surface receptors or antigens. The antisense nucleic acid molecules can also be delivered to cells using the vectors described herein. To achieve sufficient intracellular concentrations of antisense molecules, vector constructs in which the antisense nucleic acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an α-anomeric nucleic acid molecule. An α-anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual β-units, the strands run parallel to each other (Gaultier *et al.* (1987) *Nucleic Acids Res* 15: 6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (Inoue *et al.* (1987) *Nucleic Acids Res* 15: 6131-6148) or a chimeric RNA -DNA analogue (Inoue *et al.* (1987) *FEBS Lett* 215: 327-330).

4.4 RIBOZYMES AND PNA MOIETIES

In still another embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity that are capable of cleaving a single-stranded nucleic acid, such as a mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) Nature 334:585-591)) can be used to catalytically cleave a mRNA transcripts to thereby inhibit translation of a mRNA. A ribozyme having specificity for a nucleic acid of the invention can be

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designed based upon the nucleotide sequence of a DNA disclosed herein (*i.e.*, SEQ ID NO:1-1009). For example, a derivative of a Tetrahymena L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in a SECX-encoding mRNA. See, *e.g.*, Cech *et al.* U.S. Pat. No. 4,987,071; and Cech *et al.* U.S. Pat. No. 5,116,742. Alternatively, SECX mRNA can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. See, *e.g.*, Bartel *et al.*, (1993) *Science* 261:1411-1418.

Alternatively, gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region (e.g., promoter and/or enhancers) to form triple helical structures that prevent transcription of the gene in target cells. See generally, Helene. (1991) Anticancer Drug Des. 6: 569-84; Helene. et al. (1992) Ann. N.Y. Acad. Sci. 660:27-36; and Maher (1992) Bioassays 14: 807-15.

In various embodiments, the nucleic acids of the invention can be modified at the base moiety, sugar moiety or phosphate backbone to improve, e.g., the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids (see Hyrup et al. (1996) Bioorg Med Chem 4: 5-23). As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics, e.g., DNA mimics, in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained. The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength. The synthesis of PNA oligomers can be performed using standard solid phase peptide synthesis protocols as described in Hyrup et al. (1996) above; Perry-O'Keefe et al. (1996) PNAS 93: 14670-675.

PNAs of the invention can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, *e.g.*, inducing transcription or translation arrest or inhibiting replication. PNAs of the invention can also be used, *e.g.*, in the analysis of single base pair mutations in a gene by, *e.g.*, PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, *e.g.*, S1 nucleases (Hyrup B. (1996) above); or as probes or primers for DNA sequence and hybridization (Hyrup *et al.* (1996), above; Perry-O'Keefe (1996), above).

In another embodiment, PNAs of the invention can be modified, e.g., to enhance their stability or cellular uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras can be generated that may

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combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes, *e.g.*, RNase H and DNA polymerases, to interact with the DNA portion while the PNA portion would provide high binding affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths selected in terms of base stacking, number of bonds between the nucleobases, and orientation (Hyrup (1996) above). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup (1996) above and Finn *et al.* (1996) *Nucl Acids Res* 24: 3357-63. For example, a DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry, and modified nucleoside analogs, *e.g.*, 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine phosphoramidite, can be used between the PNA and the 5' end of DNA (Mag *et al.* (1989) *Nucl Acid Res* 17: 5973-88). PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment (Finn *et al.* (1996) above). Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment. See, Petersen *et al.* (1975) *Bioorg Med Chem Lett* 5: 1119-11124.

In other embodiments, the oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors in vivo), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. W088/09810) or the blood-brain barrier (see, e.g., PCT Publication No. W089/10134). In addition, oligonucleotides can be modified with hybridization triggered cleavage agents (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5: 539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, a hybridization triggered cross-linking agent, a transport agent, a hybridization-triggered cleavage agent, etc.

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4.5 HOSTS

The present invention further provides host cells genetically engineered to contain the polynucleotides of the invention. For example, such host cells may contain nucleic acids of the invention introduced into the host cell using known transformation, transfection or infection methods. The present invention still further provides host cells genetically engineered to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell.

Knowledge of nucleic acid sequences allows for modification of cells to permit, or increase, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous

recombination) to provide increased polypeptide expression by replacing, in whole or in part, the . naturally occurring promoter with all or part of a heterologous promoter so that the cells express the polypeptide at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the encoding sequences. See, for example, PCT International Publication No. WO94/12650, PCT International Publication No. WO92/20808, and PCT International Publication No. WO91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the coding sequence, amplification of the marker DNA by standard selection methods results in coamplification of the desired protein coding sequences in the cells.

The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Introduction of the recombinant construct into the host cell can be effected by calcium phosphate transfection, DEAE, dextran mediated transfection, or electroporation (Davis, L. et al., *Basic Methods in Molecular Biology* (1986)). The host cells containing one of the polynucleotides of the invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, Cv-1 cell, COS cells, 293 cells, and Sf9 cells, as well as prokaryotic host such as *E. coli* and *B. subtilis*. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level. Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by Sambrook, et al., in Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman, Cell 23:175 (1981). Other cell lines capable of expressing a compatible vector are, for example, the C127, monkey COS cells, Chinese Hamster Ovary (CHO) cells, human kidney 293 cells, human epidermal A431 cells, human Colo205 cells, 3T3

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cells, CV-1 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from *in vitro* culture of primary tissue, primary explants, HeLa cells, mouse L cells, BHK, HL-60, U937, HaK or Jurkat cells. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5′ flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements. Recombinant polypeptides and proteins produced in bacterial culture are usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents.

Alternatively, it may be possible to produce the protein in lower eukaryotes such as yeast or insects or in prokaryotes such as bacteria. Potentially suitable yeast strains include Saccharomyces cerevisiae, Schizosaccharomyces pombe, Kluyveromyces strains, Candida, or any yeast strain capable of expressing heterologous proteins. Potentially suitable bacterial strains include Escherichia coli, Bacillus subtilis, Salmonella typhimurium, or any bacterial strain capable of expressing heterologous proteins. If the protein is made in yeast or bacteria, it may be necessary to modify the protein produced therein, for example by phosphorylation or glycosylation of the appropriate sites, in order to obtain the functional protein. Such covalent attachments may be accomplished using known chemical or enzymatic methods.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequence include polyadenylation signals, mRNA stability elements, splice

sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the host cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.6 POLYPEPTIDES OF THE INVENTION

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising: the amino acid sequences set forth as any one of SEQ ID NO:1010-2018 or an amino acid sequence encoded by any one of the nucleotide sequences SEQ ID NO:1-1009 or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides preferably with biological or immunological activity that are encoded by: (a) a polynucleotide having any one of the nucleotide sequences set forth in SEQ ID NO:1-1009 or (b)

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polynucleotides encoding any one of the amino acid sequences set forth as SEQ ID NO:1010-2018 or (c) polynucleotides that hybridize to the complement of the polynucleotides of either (a) or (b) under stringent hybridization conditions. The invention also provides biologically active or immunologically active variants of any of the amino acid sequences set forth as SEQ ID NO:1010-2018 or the corresponding full length or mature protein; and "substantial equivalents" thereof (e.g., with at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, 86%, 87%, 88%, 89%, at least about 90%, 91%, 92%, 93%, 94%, typically at least about 95%, 96%, 97%, more typically at least about 98%, or most typically at least about 99% amino acid identity) that retain biological activity. Polypeptides encoded by allelic variants may have a similar, increased, or decreased activity compared to polypeptides comprising SEQ ID NO:1010-2018.

Fragments of the proteins of the present invention which are capable of exhibiting biological activity are also encompassed by the present invention. Fragments of the protein may be in linear form or they may be cyclized using known methods, for example, as described in H. U. Saragovi, et al., Bio/Technology 10, 773-778 (1992) and in R. S. McDowell, et al., J. Amer. Chem. Soc. 114, 9245-9253 (1992), both of which are incorporated herein by reference. Such fragments may be fused to carrier molecules such as immunoglobulins for many purposes, including increasing the valency of protein binding sites.

The present invention also provides both full-length and mature forms (for example, without a signal sequence or precursor sequence) of the disclosed proteins. The protein coding sequence is identified in the sequence listing by translation of the disclosed nucleotide sequences. The mature form of such protein may be obtained by expression of a full-length polynucleotide in a suitable mammalian cell or other host cell. The sequence of the mature form of the protein is also determinable from the amino acid sequence of the full-length form. Where proteins of the present invention are membrane bound, soluble forms of the proteins are also provided. In such forms, part or all of the regions causing the proteins to be membrane bound are deleted so that the proteins are fully secreted from the cell in which they are expressed.

Protein compositions of the present invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (e.g., an ORF) by nucleotide sequence but, due to the degeneracy of the genetic code, encode an identical polypeptide sequence. Preferred nucleic acid fragments of the present invention are the ORFs that encode proteins.

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A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. The synthetically-constructed protein sequences, by virtue of sharing primary, secondary or tertiary structural and/or conformational characteristics with proteins may possess biological properties in common therewith, including protein activity. This technique is particularly useful in producing small peptides and fragments of larger polypeptides. Fragments are useful, for example, in generating antibodies against the native polypeptide. Thus, they may be employed as biologically active or immunological substitutes for natural, purified proteins in screening of therapeutic compounds and in immunological processes for the development of antibodies.

The polypeptides and proteins of the present invention can alternatively be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. One skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The invention also relates to methods for producing a polypeptide comprising growing a culture of host cells of the invention in a suitable culture medium, and purifying the protein from the cells or the culture in which the cells are grown. For example, the methods of the invention include a process for producing a polypeptide in which a host cell containing a suitable expression vector that includes a polynucleotide of the invention is cultured under conditions that allow expression of the encoded polypeptide. The polypeptide can be recovered from the culture, conveniently from the culture medium, or from a lysate prepared from the host cells and further purified. Preferred embodiments include those in which the protein produced by such process is a full length or mature form of the protein.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily follow known methods for isolating polypeptides and proteins in order to obtain one of the isolated polypeptides or proteins of the present invention. These include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography. See, e.g., Scopes, Protein Purification: Principles and Practice, Springer-Verlag (1994); Sambrook, et al., in Molecular Cloning: A Laboratory

Manual; Ausubel et al., Current Protocols in Molecular Biology. Polypeptide fragments that

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retain biological/immunological activity include fragments comprising greater than about 100 amino acids, or greater than about 200 amino acids, and fragments that encode specific protein domains.

The purified polypeptides can be used in *in vitro* binding assays which are well known in the art to identify molecules which bind to the polypeptides. These molecules include but are not limited to, for *e.g.*, small molecules, molecules from combinatorial libraries, antibodies or other proteins. The molecules identified in the binding assay are then tested for antagonist or agonist activity in *in vivo* tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

In addition, the peptides of the invention or molecules capable of binding to the peptides may be complexed with toxins, *e.g.*, ricin or cholera, or with other compounds that are toxic to cells. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for SEQ ID NO:1010-2018.

The protein of the invention may also be expressed as a product of transgenic animals, e.g., as a component of the milk of transgenic cows, goats, pigs, or sheep which are characterized by somatic or germ cells containing a nucleotide sequence encoding the protein.

The proteins provided herein also include proteins characterized by amino acid sequences similar to those of purified proteins but into which modification are naturally provided or deliberately engineered. For example, modifications, in the peptide or DNA sequence, can be made by those skilled in the art using known techniques. Modifications of interest in the protein sequences may include the alteration, substitution, replacement, insertion or deletion of a selected amino acid residue in the coding sequence. For example, one or more of the cysteine residues may be deleted or replaced with another amino acid to alter the conformation of the molecule. Techniques for such alteration, substitution, replacement, insertion or deletion are well known to those skilled in the art (see, e.g., U.S. Pat. No. 4,518,584). Preferably, such alteration, substitution, replacement, insertion or deletion retains the desired activity of the protein. Regions of the protein that are important for the protein function can be determined by various methods known in the art including the alanine-scanning method which involved systematic substitution of single or strings of amino acids with alanine, followed by testing the resulting alanine-containing variant for biological activity. This type of analysis determines the importance of the substituted amino acid(s) in biological activity. Regions of the protein that are important for protein function may be determined by the eMATRIX program.

Other fragments and derivatives of the sequences of proteins which would be expected to retain protein activity in whole or in part and are useful for screening or other immunological

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methodologies may also be easily made by those skilled in the art given the disclosures herein. Such modifications are encompassed by the present invention.

The protein may also be produced by operably linking the isolated polynucleotide of the invention to suitable control sequences in one or more insect expression vectors, and employing an insect expression system. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, *e.g.*, Invitrogen, San Diego, Calif., U.S.A. (the MaxBatTM kit), and such methods are well known in the art, as described in Summers and Smith, Texas Agricultural Experiment Station Bulletin No. 1555 (1987), incorporated herein by reference. As used herein, an insect cell capable of expressing a polynucleotide of the present invention is "transformed."

The protein of the invention may be prepared by culturing transformed host cells under culture conditions suitable to express the recombinant protein. The resulting expressed protein may then be purified from such culture (*i.e.*, from culture medium or cell extracts) using known purification processes, such as gel filtration and ion exchange chromatography. The purification of the protein may also include an affinity column containing agents which will bind to the protein; one or more column steps over such affinity resins as concanavalin A-agarose, heparin-toyopearlTM or Cibacrom blue 3GA SepharoseTM; one or more steps involving hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; or immunoaffinity chromatography.

Alternatively, the protein of the invention may also be expressed in a form which will facilitate purification. For example, it may be expressed as a fusion protein, such as those of maltose binding protein (MBP), glutathione-S-transferase (GST) or thioredoxin (TRX), or as a His tag. Kits for expression and purification of such fusion proteins are commercially available from New England BioLab (Beverly, Mass.), Pharmacia (Piscataway, N.J.) and Invitrogen, respectively. The protein can also be tagged with an epitope and subsequently purified by using a specific antibody directed to such epitope. One such epitope ("FLAG®") is commercially available from Kodak (New Haven, Conn.).

Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the protein. Some or all of the foregoing purification steps, in various combinations, can also be employed to provide a substantially homogeneous isolated recombinant protein. The protein thus purified is substantially free of other mammalian proteins and is defined in accordance with the present invention as an "isolated protein."

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The polypeptides of the invention include analogs (variants). This embraces fragments, as well as peptides in which one or more amino acids has been deleted, inserted, or substituted. Also, analogs of the polypeptides of the invention embrace fusions of the polypeptides or modifications of the polypeptides of the invention, wherein the polypeptide or analog is fused to another moiety or moieties, *e.g.*, targeting moiety or another therapeutic agent. Such analogs may exhibit improved properties such as activity and/or stability. Examples of moieties which may be fused to the polypeptide or an analog include, for example, targeting moieties which provide for the delivery of polypeptide to pancreatic cells, *e.g.*, antibodies to pancreatic cells, antibodies to immune cells such as T-cells, monocytes, dendritic cells, granulocytes, etc., as well as receptor and ligands expressed on pancreatic or immune cells. Other moieties which may be fused to the polypeptide include therapeutic agents which are used for treatment, for example, immunosuppressive drugs such as cyclosporin, SK506, azathioprine, CD3 antibodies and steroids. Also, polypeptides may be fused to immune modulators, and other cytokines such as alpha or beta interferon.

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4.6.1 DETERMINING POLYPEPTIDE AND POLYNUCLEOTIDE IDENTITY AND SIMILARITY

Preferred identity and/or similarity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in computer 20 programs including, but are not limited to, the GCG program package, including GAP (Devereux, J., et al., Nucleic Acids Research 12(1):387 (1984); Genetics Computer Group, University of Wisconsin, Madison, WI), BLASTP, BLASTN, BLASTX, FASTA (Altschul, S.F. et al., J. Molec. Biol. 215:403-410 (1990), PSI-BLAST (Altschul S.F. et al., Nucleic Acids Res. vol. 25, pp. 3389-3402, herein incorporated by reference), eMatrix software (Wu et al., J. Comp. 25 Biol., Vol. 6, pp. 219-235 (1999), herein incorporated by reference), eMotif software (Nevill-Manning et al, ISMB-97, Vol. 4, pp. 202-209, herein incorporated by reference), pFam software (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1), pp. 320-322 (1998), herein incorporated by reference) and the Kyte-Doolittle hydrophobocity prediction algorithm (J. Mol Biol, 157, pp. 105-31 (1982), incorporated herein by reference). The BLAST programs are publicly available 30 from the National Center for Biotechnology Information (NCBI) and other sources (BLAST Manual, Altschul, S., et al. NCB NLM NIH Bethesda, MD 20894; Altschul, S., et al., J. Mol. Biol. 215:403-410 (1990).

4.7 CHIMERIC AND FUSION PROTEINS

The invention also provides chimeric or fusion proteins. As used herein, a "chimeric protein" or "fusion protein" comprises a polypeptide of the invention operatively linked to

another polypeptide. Within a fusion protein the polypeptide according to the invention can correspond to all or a portion of a protein according to the invention. In one embodiment, a fusion protein comprises at least one biologically active portion of a protein according to the invention. In another embodiment, a fusion protein comprises at least two biologically active portions of a protein according to the invention. Within the fusion protein, the term "operatively linked" is intended to indicate that the polypeptide according to the invention and the other polypeptide are fused in-frame to each other. The polypeptide can be fused to the N-terminus or C-terminus.

For example, in one embodiment a fusion protein comprises a polypeptide according to the invention operably linked to the extracellular domain of a second protein.

In another embodiment, the fusion protein is a GST-fusion protein in which the polypeptide sequences of the invention are fused to the C-terminus of the GST (i.e., glutathione S-transferase) sequences.

In another embodiment, the fusion protein is an immunoglobulin fusion protein in which the polypeptide sequences according to the invention comprises one or more domains are fused to sequences derived from a member of the immunoglobulin protein family. The immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between a ligand and a protein of the invention on the surface of a cell, to thereby suppress signal transduction *in vivo*. The immunoglobulin fusion proteins can be used to affect the bioavailability of a cognate ligand. Inhibition of the ligand/protein interaction may be useful therapeutically for both the treatment of proliferative and differentiative disorders, *e,g.*, cancer as well as modulating (*e.g.*, promoting or inhibiting) cell survival. Moreover, the immunoglobulin fusion proteins of the invention can be used as immunogens to produce antibodies in a subject, to purify ligands, and in screening assays to identify molecules that inhibit the interaction of a polypeptide of the invention with a ligand.

A chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, *e.g.*, by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for

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example, Ausubel et al. (eds.) Current Protocols in Molecular Biology, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). A nucleic acid encoding a polypeptide of the invention can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the protein of the invention.

4.8 GENE THERAPY

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Mutations in the polynucleotides of the invention gene may result in loss of normal function of the encoded protein. The invention thus provides gene therapy to restore normal activity of the polypeptides of the invention; or to treat disease states involving polypeptides of the invention. Delivery of a functional gene encoding polypeptides of the invention to appropriate cells is effected ex vivo, in situ, or in vivo by use of vectors, and more particularly viral vectors (e.g., adenovirus, adeno-associated virus, or a retrovirus), or ex vivo by use of physical DNA transfer methods (e.g., liposomes or chemical treatments). See, for example, Anderson, Nature, supplement to vol. 392, no. 6679, pp.25-20 (1998). For additional reviews of gene therapy technology see Friedmann, Science, 244: 1275-1281 (1989); Verma, Scientific American: 68-84 (1990); and Miller, Nature, 357: 455-460 (1992). Introduction of any one of the nucleotides of the present invention or a gene encoding the polypeptides of the present invention can also be accomplished with extrachromosomal substrates (transient expression) or artificial chromosomes (stable expression). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes. Alternatively, it is contemplated that in other human disease states, preventing the expression of or inhibiting the activity of polypeptides of the invention will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate the expression of polypeptides of the invention.

Other methods inhibiting expression of a protein include the introduction of antisense molecules to the nucleic acids of the present invention, their complements, or their translated RNA sequences, by methods known in the art. Further, the polypeptides of the present invention can be inhibited by using targeted deletion methods, or the insertion of a negative regulatory element such as a silencer, which is tissue specific.

The present invention still further provides cells genetically engineered *in vivo* to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in

the cell. These methods can be used to increase or decrease the expression of the polynucleotides of . the present invention.

Knowledge of DNA sequences provided by the invention allows for modification of cells to permit, increase, or decrease, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the protein at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the desired protein encoding sequences. See, for example, PCT International Publication No. WO 94/12650, PCT International Publication No. WO 92/20808, and PCT International Publication No. WO 91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the desired protein coding sequence, amplification of the marker DNA by standard selection methods results in co-amplification of the desired protein coding sequences in the cells.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequences include polyadenylation signals, mRNA stability elements, splice sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are

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added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

4.9 TRANSGENIC ANIMALS

In preferred methods to determine biological functions of the polypeptides of the invention in vivo, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of a promoter of the polynucleotides of the invention is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous

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promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

The polynucleotides of the present invention also make possible the development, through, e.g., homologous recombination or knock out strategies, of animals that fail to express polypeptides of the invention or that express a variant polypeptide. Such animals are useful as models for studying the *in vivo* activities of polypeptide as well as for studying modulators of the polypeptides of the invention.

In preferred methods to determine biological functions of the polypeptides of the invention *in vivo*, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of the polynucleotides of the invention promoter is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

4.10 USES AND BIOLOGICAL ACTIVITY

The polynucleotides and proteins of the present invention are expected to exhibit one or more of the uses or biological activities (including those associated with assays cited herein) identified herein. Uses or activities described for proteins of the present invention may be provided by administration or use of such proteins or of polynucleotides encoding such proteins (such as, for example, in gene therapies or vectors suitable for introduction of DNA). The mechanism underlying the particular condition or pathology will dictate whether the

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polypeptides of the invention, the polynucleotides of the invention or modulators (activators or inhibitors) thereof would be beneficial to the subject in need of treatment. Thus, "therapeutic compositions of the invention" include compositions comprising isolated polynucleotides (including recombinant DNA molecules, cloned genes and degenerate variants thereof) or polypeptides of the invention (including full length protein, mature protein and truncations or domains thereof), or compounds and other substances that modulate the overall activity of the target gene products, either at the level of target gene/protein expression or target protein activity. Such modulators include polypeptides, analogs, (variants), including fragments and fusion proteins, antibodies and other binding proteins; chemical compounds that directly or indirectly activate or inhibit the polypeptides of the invention (identified, *e.g.*, via drug screening assays as described herein); antisense polynucleotides and polynucleotides suitable for triple helix formation; and in particular antibodies or other binding partners that specifically recognize one or more epitopes of the polypeptides of the invention.

The polypeptides of the present invention may likewise be involved in cellular activation or in one of the other physiological pathways described herein.

4.10.1 RESEARCH USES AND UTILITIES

The polynucleotides provided by the present invention can be used by the research community for various purposes. The polynucleotides can be used to express recombinant protein for analysis, characterization or therapeutic use; as markers for tissues in which the corresponding protein is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in disease states); as molecular weight markers on gels; as chromosome markers or tags (when labeled) to identify chromosomes or to map related gene positions; to compare with endogenous DNA sequences in patients to identify potential genetic disorders; as probes to hybridize and thus discover novel, related DNA sequences; as a source of information to derive PCR primers for genetic fingerprinting; as a probe to "subtract-out" known sequences in the process of discovering other novel polynucleotides; for selecting and making oligomers for attachment to a "gene chip" or other support, including for examination of expression patterns; to raise anti-protein antibodies using DNA immunization techniques; and as an antigen to raise anti-DNA antibodies or elicit another immune response. Where the polynucleotide encodes a protein which binds or potentially binds to another protein (such as, for example, in a receptor-ligand interaction), the polynucleotide can also be used in interaction trap assays (such as, for example, that described in Gyuris et al., Cell 75:791-803 (1993)) to identify polynucleotides encoding the other protein with which binding occurs or to identify inhibitors of the binding interaction.

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The polypeptides provided by the present invention can similarly be used in assays to determine biological activity, including in a panel of multiple proteins for high-throughput screening; to raise antibodies or to elicit another immune response; as a reagent (including the labeled reagent) in assays designed to quantitatively determine levels of the protein (or its receptor) in biological fluids; as markers for tissues in which the corresponding polypeptide is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in a disease state); and, of course, to isolate correlative receptors or ligands. Proteins involved in these binding interactions can also be used to screen for peptide or small molecule inhibitors or agonists of the binding interaction.

Any or all of these research utilities are capable of being developed into reagent grade or kit format for commercialization as research products.

Methods for performing the uses listed above are well known to those skilled in the art. References disclosing such methods include without limitation "Molecular Cloning: A Laboratory Manual", 2d ed., Cold Spring Harbor Laboratory Press, Sambrook, J., E. F. Fritsch and T. Maniatis eds., 1989, and "Methods in Enzymology: Guide to Molecular Cloning Techniques", Academic Press, Berger, S. L. and A. R. Kimmel eds., 1987.

4.10.2 NUTRITIONAL USES

Polynucleotides and polypeptides of the present invention can also be used as nutritional sources or supplements. Such uses include without limitation use as a protein or amino acid supplement, use as a carbon source, use as a nitrogen source and use as a source of carbohydrate. In such cases the polypeptide or polynucleotide of the invention can be added to the feed of a particular organism or can be administered as a separate solid or liquid preparation, such as in the form of powder, pills, solutions, suspensions or capsules. In the case of microorganisms, the polypeptide or polynucleotide of the invention can be added to the medium in or on which the microorganism is cultured.

4.10.3 CYTOKINE AND CELL PROLIFERATION/DIFFERENTIATION ACTIVITY

A polypeptide of the present invention may exhibit activity relating to cytokine, cell proliferation (either inducing or inhibiting) or cell differentiation (either inducing or inhibiting) activity or may induce production of other cytokines in certain cell populations. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Many protein factors discovered to date, including all known cytokines, have exhibited activity in one or more factor-dependent cell proliferation assays, and hence the assays serve as a convenient

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confirmation of cytokine activity. The activity of therapeutic compositions of the present invention is evidenced by any one of a number of routine factor dependent cell proliferation assays for cell lines including, without limitation, 32D, DA2, DA1G, T10, B9, B9/11, BaF3, MC9/G, M+(preB M+), 2E8, RB5, DA1, 123, T1165, HT2, CTLL2, TF-1, Mo7e, CMK, HUVEC, and Caco. Therapeutic compositions of the invention can be used in the following:

Assays for T-cell or thymocyte proliferation include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Bertagnolli et al., J. Immunol. 145:1706-1712, 1990; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Bertagnolli, et al., I. Immunol. 149:3778-3783, 1992; Bowman et al., I: Immunol. 152:1756-1761, 1994.

Assays for cytokine production and/or proliferation of spleen cells, lymph node cells or thymocytes include, without limitation, those described in: Polyclonal T cell stimulation, Kruisbeek, A. M. and Shevach, E. M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.12.1-3.12.14, John Wiley and Sons, Toronto. 1994; and Measurement of mouse and human interleukin-γ, Schreiber, R. D. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.8.1-6.8.8, John Wiley and Sons, Toronto. 1994.

Assays for proliferation and differentiation of hematopoietic and lymphopoietic cells 20 include, without limitation, those described in: Measurement of Human and Murine Interleukin 2 and Interleukin 4, Bottomly, K., Davis, L. S. and Lipsky, P. E. In Current Protocols in Immunology, J. E. e.a. Coligan eds. Vol 1 pp. 6.3.1-6.3.12, John Wiley and Sons, Toronto. 1991: deVries et al., J. Exp. Med. 173:1205-1211, 1991; Moreau et al., Nature 336:690-692, 1988; Greenberger et al., Proc. Natl. Acad. Sci. U.S.A. 80:2931-2938, 1983; Measurement of mouse 25 and human interleukin 6--Nordan, R. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.6.1-6.6.5, John Wiley and Sons, Toronto. 1991; Smith et al., Proc. Natl. Aced. Sci. U.S.A. 83:1857-1861, 1986; Measurement of human Interleukin 11--Bennett, F., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology, J. E. Coligan eds. Vol 1 pp. 6.15.1 John Wiley and Sons, Toronto. 1991; Measurement of mouse and human Interleukin 30 9--Ciarletta, A., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.13.1, John Wiley and Sons, Toronto. 1991.

Assays for T-cell clone responses to antigens (which will identify, among others, proteins that affect APC-T cell interactions as well as direct T-cell effects by measuring proliferation and cytokine production) include, without limitation, those described in: Current Protocols in

Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober,

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Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse. Lymphocyte Function; Chapter 6, Cytokines and their cellular receptors; Chapter 7, Immunologic studies in Humans); Weinberger et al., Proc. Natl. Acad. Sci. USA 77:6091-6095, 1980; Weinberger et al., Eur. J. Immun. 11:405-411, 1981; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988.

4.10.4 STEM CELL GROWTH FACTOR ACTIVITY

A polypeptide of the present invention may exhibit stem cell growth factor activity and be involved in the proliferation, differentiation and survival of pluripotent and totipotent stem cells including primordial germ cells, embryonic stem cells, hematopoietic stem cells and/or germ line stem cells. Administration of the polypeptide of the invention to stem cells *in vivo* or *ex vivo* is expected to maintain and expand cell populations in a totipotential or pluripotential state which would be useful for re-engineering damaged or diseased tissues, transplantation, manufacture of bio-pharmaceuticals and the development of bio-sensors. The ability to produce large quantities of human cells has important working applications for the production of human proteins which currently must be obtained from non-human sources or donors, implantation of cells to treat diseases such as Parkinson's, Alzheimer's and other neurodegenerative diseases; tissues for grafting such as bone marrow, skin, cartilage, tendons, bone, muscle (including cardiac muscle), blood vessels, cornea, neural cells, gastrointestinal cells and others; and organs for transplantation such as kidney, liver, pancreas (including islet cells), heart and lung.

It is contemplated that multiple different exogenous growth factors and/or cytokines may be administered in combination with the polypeptide of the invention to achieve the desired effect, including any of the growth factors listed herein, other stem cell maintenance factors, and specifically including stem cell factor (SCF), leukemia inhibitory factor (LIF), Flt-3 ligand (Flt-3L), any of the interleukins, recombinant soluble IL-6 receptor fused to IL-6, macrophage inflammatory protein 1-alpha (MIP-1-alpha), G-CSF, GM-CSF, thrombopoietin (TPO), platelet factor 4 (PF-4), platelet-derived growth factor (PDGF), neural growth factors and basic fibroblast growth factor (bFGF).

Since totipotent stem cells can give rise to virtually any mature cell type, expansion of these cells in culture will facilitate the production of large quantities of mature cells. Techniques for culturing stem cells are known in the art and administration of polypeptides of the invention, optionally with other growth factors and/or cytokines, is expected to enhance the survival and proliferation of the stem cell populations. This can be accomplished by direct administration of the polypeptide of the invention to the culture medium. Alternatively, stroma cells transfected with a polynucleotide that encodes for the polypeptide of the invention can be used as a feeder

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layer for the stem cell populations in culture or in vivo. Stromal support cells for feeder layers may include embryonic bone marrow fibroblasts, bone marrow stromal cells, fetal liver cells, or cultured embryonic fibroblasts (see U.S. Patent No. 5,690,926).

Stem cells themselves can be transfected with a polynucleotide of the invention to induce autocrine expression of the polypeptide of the invention. This will allow for generation of undifferentiated totipotential/pluripotential stem cell lines that are useful as is or that can then be differentiated into the desired mature cell types. These stable cell lines can also serve as a source of undifferentiated totipotential/pluripotential mRNA to create cDNA libraries and templates for polymerase chain reaction experiments. These studies would allow for the isolation and identification of differentially expressed genes in stem cell populations that regulate stem cell proliferation and/or maintenance.

Expansion and maintenance of totipotent stem cell populations will be useful in the treatment of many pathological conditions. For example, polypeptides of the present invention may be used to manipulate stem cells in culture to give rise to neuroepithelial cells that can be used to augment or replace cells damaged by illness, autoimmune disease, accidental damage or genetic disorders. The polypeptide of the invention may be useful for inducing the proliferation of neural cells and for the regeneration of nerve and brain tissue, *i.e.* for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders which involve degeneration, death or trauma to neural cells or nerve tissue. In addition, the expanded stem cell populations can also be genetically altered for gene therapy purposes and to decrease host rejection of replacement tissues after grafting or implantation.

Expression of the polypeptide of the invention and its effect on stem cells can also be manipulated to achieve controlled differentiation of the stem cells into more differentiated cell types. A broadly applicable method of obtaining pure populations of a specific differentiated cell type from undifferentiated stem cell populations involves the use of a cell-type specific promoter driving a selectable marker. The selectable marker allows only cells of the desired type to survive. For example, stem cells can be induced to differentiate into cardiomyocytes (Wobus et al., Differentiation, 48: 173-182, (1991); Klug et al., J. Clin. Invest., 98(1): 216-224, (1998)) or skeletal muscle cells (Browder, L. W. In: *Principles of Tissue Engineering eds.* Lanza et al., Academic Press (1997)). Alternatively, directed differentiation of stem cells can be accomplished by culturing the stem cells in the presence of a differentiation factor such as retinoic acid and an antagonist of the polypeptide of the invention which would inhibit the effects of endogenous stem cell factor activity and allow differentiation to proceed.

In vitro cultures of stem cells can be used to determine if the polypeptide of the invention exhibits stem cell growth factor activity. Stem cells are isolated from any one of various cell

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sources (including hematopoietic stem cells and embryonic stem cells) and cultured on a feeder layer, as described by Thompson et al. Proc. Natl. Acad. Sci, U.S.A., 92: 7844-7848 (1995), in the presence of the polypeptide of the invention alone or in combination with other growth factors or cytokines. The ability of the polypeptide of the invention to induce stem cells proliferation is determined by colony formation on semi-solid support *e.g.* as described by Bernstein et al., Blood, 77: 2316-2321 (1991).

4.10.5 HEMATOPOIESIS REGULATING ACTIVITY

A polypeptide of the present invention may be involved in regulation of hematopoiesis and, consequently, in the treatment of myeloid or lymphoid cell disorders. Even marginal biological activity in support of colony forming cells or of factor-dependent cell lines indicates involvement in regulating hematopoiesis, e.g. in supporting the growth and proliferation of erythroid progenitor cells alone or in combination with other cytokines, thereby indicating utility, for example, in treating various anemias or for use in conjunction with irradiation/chemotherapy to stimulate the production of erythroid precursors and/or erythroid cells; in supporting the growth and proliferation of myeloid cells such as granulocytes and monocytes/macrophages (i.e., traditional CSF activity) useful, for example, in conjunction with chemotherapy to prevent or treat consequent myelo-suppression; in supporting the growth and proliferation of megakaryocytes and consequently of platelets thereby allowing prevention or treatment of various platelet disorders such as thrombocytopenia, and generally for use in place of or complimentary to platelet transfusions; and/or in supporting the growth and proliferation of hematopoietic stem cells which are capable of maturing to any and all of the above-mentioned hematopoietic cells and therefore find therapeutic utility in various stem cell disorders (such as those usually treated with transplantation, including, without limitation, aplastic anemia and paroxysmal nocturnal hemoglobinuria), as well as in repopulating the stem cell compartment post irradiation/chemotherapy, either in-vivo or ex-vivo (i.e., in conjunction with bone marrow transplantation or with peripheral progenitor cell transplantation (homologous or heterologous)) as normal cells or genetically manipulated for gene therapy.

Therapeutic compositions of the invention can be used in the following:

Suitable assays for proliferation and differentiation of various hematopoietic lines are cited above.

Assays for embryonic stem cell differentiation (which will identify, among others, proteins that influence embryonic differentiation hematopoiesis) include, without limitation, those described in: Johansson et al. Cellular Biology 15:141-151, 1995; Keller et al., Molecular and Cellular Biology 13:473-486, 1993; McClanahan et al., Blood 81:2903-2915, 1993.

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Assays for stem cell survival and differentiation (which will identify, among others, proteins that regulate lympho-hematopoiesis) include, without limitation, those described in: Methylcellulose colony forming assays, Freshney, M. G. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 265-268, Wiley-Liss, Inc., New York, N.Y. 1994; Hirayama et al., Proc. Natl. Acad. Sci. USA 89:5907-5911, 1992; Primitive hematopoietic colony forming cells with high proliferative potential, McNiece, I. K. and Briddell, R. A. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 23-39, Wiley-Liss, Inc., New York, N.Y. 1994; Neben et al., Experimental Hematology 22:353-359, 1994; Cobblestone area forming cell assay, Ploemacher, R. E. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 1-21, Wiley-Liss, Inc., New York, N.Y. 1994; Long term bone marrow cultures in the presence of stromal cells, Spooncer, E., Dexter, M. and Allen, T. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 163-179, Wiley-Liss, Inc., New York, N.Y. 1994; Long term culture initiating cell assay, Sutherland, H. J. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 139-162, Wiley-Liss, Inc., New York, N.Y. 1994.

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4.10.6 TISSUE GROWTH ACTIVITY

A polypeptide of the present invention also may be involved in bone, cartilage, tendon, ligament and/or nerve tissue growth or regeneration, as well as in wound healing and tissue repair and replacement, and in healing of burns, incisions and ulcers.

A polypeptide of the present invention which induces cartilage and/or bone growth in circumstances where bone is not normally formed, has application in the healing of bone fractures and cartilage damage or defects in humans and other animals. Compositions of a polypeptide, antibody, binding partner, or other modulator of the invention may have prophylactic use in closed as well as open fracture reduction and also in the improved fixation of artificial joints. De novo bone formation induced by an osteogenic agent contributes to the repair of congenital, trauma induced, or oncologic resection induced craniofacial defects, and also is useful in cosmetic plastic surgery.

A polypeptide of this invention may also be involved in attracting bone-forming cells, stimulating growth of bone-forming cells, or inducing differentiation of progenitors of bone-forming cells. Treatment of osteoporosis, osteoarthritis, bone degenerative disorders, or periodontal disease, such as through stimulation of bone and/or cartilage repair or by blocking inflammation or processes of tissue destruction (collagenase activity, osteoclast activity, etc.) mediated by inflammatory processes may also be possible using the composition of the invention.

Another category of tissue regeneration activity that may involve the polypeptide of the present invention is tendon/ligament formation. Induction of tendon/ligament-like tissue or other tissue formation in circumstances where such tissue is not normally formed, has application in the healing of tendon or ligament tears, deformities and other tendon or ligament defects in humans and other animals. Such a preparation employing a tendon/ligament-like tissue inducing protein may have prophylactic use in preventing damage to tendon or ligament tissue, as well as use in the improved fixation of tendon or ligament to bone or other tissues, and in repairing defects to tendon or ligament tissue. De novo tendon/ligament-like tissue formation induced by a composition of the present invention contributes to the repair of congenital, trauma induced, or other tendon or ligament defects of other origin, and is also useful in cosmetic plastic surgery for attachment or repair of tendons or ligaments. The compositions of the present invention may provide environment to attract tendon- or ligament-forming cells, stimulate growth of tendon- or ligament-forming cells, induce differentiation of progenitors of tendon- or ligament-forming cells, or induce growth of tendon/ligament cells or progenitors ex vivo for return in vivo to effect tissue repair. The compositions of the invention may also be useful in the treatment of tendinitis, carpal tunnel syndrome and other tendon or ligament defects. The compositions may also include an appropriate matrix and/or sequestering agent as a carrier as is well known in the art.

The compositions of the present invention may also be useful for proliferation of neural cells and for regeneration of nerve and brain tissue, *i.e.* for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders, which involve degeneration, death or trauma to neural cells or nerve tissue. More specifically, a composition may be used in the treatment of diseases of the peripheral nervous system, such as peripheral nerve injuries, peripheral neuropathy and localized neuropathies, and central nervous system diseases, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome. Further conditions which may be treated in accordance with the present invention include mechanical and traumatic disorders, such as spinal cord disorders, head trauma and cerebrovascular diseases such as stroke. Peripheral neuropathies resulting from chemotherapy or other medical therapies may also be treatable using a composition of the invention.

Compositions of the invention may also be useful to promote better or faster closure of non-healing wounds, including without limitation pressure ulcers, ulcers associated with vascular insufficiency, surgical and traumatic wounds, and the like.

Compositions of the present invention may also be involved in the generation or regeneration of other tissues, such as organs (including, for example, pancreas, liver, intestine,

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kidney, skin, endothelium), muscle (smooth, skeletal or cardiac) and vascular (including vascular endothelium) tissue, or for promoting the growth of cells comprising such tissues. Part of the desired effects may be by inhibition or modulation of fibrotic scarring may allow normal tissue to regenerate. A polypeptide of the present invention may also exhibit angiogenic activity.

A composition of the present invention may also be useful for gut protection or regeneration and treatment of lung or liver fibrosis, reperfusion injury in various tissues, and conditions resulting from systemic cytokine damage.

A composition of the present invention may also be useful for promoting or inhibiting differentiation of tissues described above from precursor tissues or cells; or for inhibiting the growth of tissues described above.

Therapeutic compositions of the invention can be used in the following:

Assays for tissue generation activity include, without limitation, those described in: International Patent Publication No. WO95/16035 (bone, cartilage, tendon); International Patent Publication No. WO95/05846 (nerve, neuronal); International Patent Publication No. WO91/07491 (skin, endothelium).

Assays for wound healing activity include, without limitation, those described in: Winter, Epidermal Wound Healing, pps. 71-112 (Maibach, H. I. and Rovee, D. T., eds.), Year Book Medical Publishers, Inc., Chicago, as modified by Eaglstein and Mertz, J. Invest. Dermatol 71:382-84 (1978).

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4.10.7 IMMUNE STIMULATING OR SUPPRESSING ACTIVITY

A polypeptide of the present invention may also exhibit immune stimulating or immune suppressing activity, including without limitation the activities for which assays are described herein. A polynucleotide of the invention can encode a polypeptide exhibiting such activities. A protein may be useful in the treatment of various immune deficiencies and disorders (including severe combined immunodeficiency (SCID)), e.g., in regulating (up or down) growth and proliferation of T and/or B lymphocytes, as well as effecting the cytolytic activity of NK cells and other cell populations. These immune deficiencies may be genetic or be caused by viral (e.g., HIV) as well as bacterial or fungal infections, or may result from autoimmune disorders. More specifically, infectious diseases causes by viral, bacterial, fungal or other infection may be treatable using a protein of the present invention, including infections by HIV, hepatitis viruses, herpes viruses, mycobacteria, Leishmania spp., malaria spp. and various fungal infections such as candidiasis. Of course, in this regard, proteins of the present invention may also be useful where a boost to the immune system generally may be desirable, i.e., in the treatment of cancer.

Autoimmune disorders which may be treated using a protein of the present invention include, for example, connective tissue disease, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary inflammation, Guillain-Barre syndrome, autoimmune thyroiditis, insulin dependent diabetes mellitis, myasthenia gravis, graft-versus-host disease and autoimmune inflammatory eye disease. Such a protein (or antagonists thereof, including antibodies) of the present invention may also to be useful in the treatment of allergic reactions and conditions (e.g., anaphylaxis, serum sickness, drug reactions, food allergies, insect venom allergies, mastocytosis, allergic rhinitis, hypersensitivity pneumonitis, urticaria, angioedema, eczema, atopic dermatitis, allergic contact dermatitis, erythema multiforme, Stevens-Johnson syndrome, allergic conjunctivitis, atopic keratoconjunctivitis, venereal keratoconjunctivitis, giant papillary conjunctivitis and contact allergies), such as asthma (particularly allergic asthma) or other respiratory problems. Other conditions, in which immune suppression is desired (including, for example, organ transplantation), may also be treatable using a protein (or antagonists thereof) of the present invention. The therapeutic effects of the polypeptides or antagonists thereof on allergic reactions can be evaluated by in vivo animals models such as the cumulative contact enhancement test (Lastborn et al., Toxicology 125: 59-66, 1998), skin prick test (Hoffmann et al., Allergy 54: 446-54, 1999), guinea pig skin sensitization test (Vohr et al., Arch. Toxocol. 73: 501-9), and murine local lymph node assay (Kimber et al., J. Toxicol. Environ. Health 53: 563-79).

Using the proteins of the invention it may also be possible to modulate immune responses, in a number of ways. Down regulation may be in the form of inhibiting or blocking an immune response already in progress or may involve preventing the induction of an immune response. The functions of activated T cells may be inhibited by suppressing T cell responses or by inducing specific tolerance in T cells, or both. Immunosuppression of T cell responses is generally an active, non-antigen-specific, process which requires continuous exposure of the T cells to the suppressive agent. Tolerance, which involves inducing non-responsiveness or anergy in T cells, is distinguishable from immunosuppression in that it is generally antigen-specific and persists after exposure to the tolerizing agent has ceased. Operationally, tolerance can be demonstrated by the lack of a T cell response upon reexposure to specific antigen in the absence of the tolerizing agent.

Down regulating or preventing one or more antigen functions (including without limitation B lymphocyte antigen functions (such as, for example, B7)), e.g., preventing high level lymphokine synthesis by activated T cells, will be useful in situations of tissue, skin and organ transplantation and in graft-versus-host disease (GVHD). For example, blockage of T cell function should result in reduced tissue destruction in tissue transplantation. Typically, in tissue

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transplants, rejection of the transplant is initiated through its recognition as foreign by T cells, followed by an immune reaction that destroys the transplant. The administration of a therapeutic composition of the invention may prevent cytokine synthesis by immune cells, such as T cells, and thus acts as an immunosuppressant. Moreover, a lack of costimulation may also be sufficient to anergize the T cells, thereby inducing tolerance in a subject. Induction of long-term tolerance by B lymphocyte antigen-blocking reagents may avoid the necessity of repeated administration of these blocking reagents. To achieve sufficient immunosuppression or tolerance in a subject, it may also be necessary to block the function of a combination of B lymphocyte antigens.

The efficacy of particular therapeutic compositions in preventing organ transplant rejection or GVHD can be assessed using animal models that are predictive of efficacy in humans. Examples of appropriate systems which can be used include allogeneic cardiac grafts in rats and xenogeneic pancreatic islet cell grafts in mice, both of which have been used to examine the immunosuppressive effects of CTLA4Ig fusion proteins in vivo as described in Lenschow et al., Science 257:789-792 (1992) and Turka et al., Proc. Natl. Acad. Sci USA, 89:11102-11105 (1992). In addition, murine models of GVHD (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 846-847) can be used to determine the effect of therapeutic compositions of the invention on the development of that disease.

Blocking antigen function may also be therapeutically useful for treating autoimmune diseases. Many autoimmune disorders are the result of inappropriate activation of T cells that are reactive against self tissue and which promote the production of cytokines and autoantibodies involved in the pathology of the diseases. Preventing the activation of autoreactive T cells may reduce or eliminate disease symptoms. Administration of reagents which block stimulation of T cells can be used to inhibit T cell activation and prevent production of autoantibodies or T cell-derived cytokines which may be involved in the disease process. Additionally, blocking reagents may induce antigen-specific tolerance of autoreactive T cells which could lead to long-term relief from the disease. The efficacy of blocking reagents in preventing or alleviating autoimmune disorders can be determined using a number of well-characterized animal models of human autoimmune diseases. Examples include murine experimental autoimmune encephalitis, systemic lupus erythmatosis in MRL/lpr/lpr mice or NZB hybrid mice, murine autoimmune collagen arthritis, diabetes mellitus in NOD mice and BB rats, and murine experimental myasthenia gravis (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 840-856).

Upregulation of an antigen function (e.g., a B lymphocyte antigen function), as a means of up regulating immune responses, may also be useful in therapy. Upregulation of immune responses may be in the form of enhancing an existing immune response or eliciting an initial

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immune response. For example, enhancing an immune response may be useful in cases of viral infection, including systemic viral diseases such as influenza, the common cold, and encephalitis.

Alternatively, anti-viral immune responses may be enhanced in an infected patient by removing T cells from the patient, costimulating the T cells in vitro with viral antigen-pulsed APCs either expressing a peptide of the present invention or together with a stimulatory form of a soluble peptide of the present invention and reintroducing the in vitro activated T cells into the patient. Another method of enhancing anti-viral immune responses would be to isolate infected cells from a patient, transfect them with a nucleic acid encoding a protein of the present invention as described herein such that the cells express all or a portion of the protein on their surface, and reintroduce the transfected cells into the patient. The infected cells would now be capable of delivering a costimulatory signal to, and thereby activate, T cells in vivo.

A polypeptide of the present invention may provide the necessary stimulation signal to T cells to induce a T cell mediated immune response against the transfected tumor cells. In addition, tumor cells which lack MHC class I or MHC class II molecules, or which fail to reexpress sufficient mounts of MHC class I or MHC class II molecules, can be transfected with nucleic acid encoding all or a portion of (e.g., a cytoplasmic-domain truncated portion) of an MHC class I alpha chain protein and an MHC class II beta chain protein to thereby express MHC class I or MHC class II proteins on the cell surface. Expression of the appropriate class I or class II MHC in conjunction with a peptide having the activity of a B lymphocyte antigen (e.g., B7-1, B7-2, B7-3) induces a T cell mediated immune response against the transfected tumor cell. Optionally, a gene encoding an antisense construct which blocks expression of an MHC class II associated protein, such as the invariant chain, can also be cotransfected with a DNA encoding a peptide having the activity of a B lymphocyte antigen to promote presentation of tumor associated antigens and induce tumor specific immunity. Thus, the induction of a T cell mediated immune response in a human subject may be sufficient to overcome tumor-specific tolerance in the subject.

The activity of a protein of the invention may, among other means, be measured by the following methods:

Suitable assays for thymocyte or splenocyte cytotoxicity include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Herrmann et al., Proc. Natl. Acad. Sci. USA 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J.

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Immunol. 135:1564-1572, 1985; Takai et al., I. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bowman et al., J. Virology 61:1992-1998; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Brown et al., J. Immunol. 153:3079-3092, 1994.

Assays for T-cell-dependent immunoglobulin responses and isotype switching (which will identify, among others, proteins that modulate T-cell dependent antibody responses and that affect Th1/Th2 profiles) include, without limitation, those described in: Maliszewski, J. Immunol. 144:3028-3033, 1990; and Assays for B cell function. In vitro antibody production, Mond, J. J. and Brunswick, M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.8.1-3.8.16, John Wiley and Sons, Toronto. 1994.

Mixed lymphocyte reaction (MLR) assays (which will identify, among others, proteins that generate predominantly Th1 and CTL responses) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnolli et al., J. Immunol. 149:3778-3783, 1992.

Dendritic cell-dependent assays (which will identify, among others, proteins expressed by dendritic cells that activate naive T-cells) include, without limitation, those described in: Guery et al., J. Immunol. 134:536-544, 1995; Inaba et al., Journal of Experimental Medicine 173:549-559, 1991; Macatonia et al., Journal of Immunology 154:5071-5079, 1995; Porgador et al., Journal of Experimental Medicine 182:255-260, 1995; Nair et al., Journal of Virology 67:4062-4069, 1993; Huang et al., Science 264:961-965, 1994; Macatonia et al., Journal of Experimental Medicine 169:1255-1264, 1989; Bhardwaj et al., Journal of Clinical Investigation 94:797-807, 1994; and Inaba et al., Journal of Experimental Medicine 172:631-640, 1990.

Assays for lymphocyte survival/apoptosis (which will identify, among others, proteins that prevent apoptosis after superantigen induction and proteins that regulate lymphocyte homeostasis) include, without limitation, those described in: Darzynkiewicz et al., Cytometry 13:795-808, 1992; Gorczyca et al., Leukemia 7:659-670, 1993; Gorczyca et al., Cancer Research 53:1945-1951, 1993; Itoh et al., Cell 66:233-243, 1991; Zacharchuk, Journal of Immunology 145:4037-4045, 1990; Zamai et al., Cytometry 14:891-897, 1993; Gorczyca et al., International Journal of Oncology 1:639-648, 1992.

Assays for proteins that influence early steps of T-cell commitment and development include, without limitation, those described in: Antica et al., Blood 84:111-117, 1994; Fine et al., Cellular Immunology 155:111-122, 1994; Galy et al., Blood 85:2770-2778, 1995; Toki et al.,

35 Proc. Nat. Acad Sci. USA 88:7548-7551, 1991.

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4.10.8 ACTIVIN/INHIBIN ACTIVITY

A polypeptide of the present invention may also exhibit activin- or inhibin-related activities. A polynucleotide of the invention may encode a polypeptide exhibiting such characteristics. Inhibins are characterized by their ability to inhibit the release of follicle stimulating hormone (FSH), while activins and are characterized by their ability to stimulate the release of follicle stimulating hormone (FSH). Thus, a polypeptide of the present invention, alone or in heterodimers with a member of the inhibin family, may be useful as a contraceptive based on the ability of inhibins to decrease fertility in female mammals and decrease spermatogenesis in male mammals. Administration of sufficient amounts of other inhibins can induce infertility in these mammals. Alternatively, the polypeptide of the invention, as a homodimer or as a heterodimer with other protein subunits of the inhibin group, may be useful as a fertility inducing therapeutic, based upon the ability of activin molecules in stimulating FSH release from cells of the anterior pituitary. See, for example, U.S. Pat. No. 4,798,885. A polypeptide of the invention may also be useful for advancement of the onset of fertility in sexually immature mammals, so as to increase the lifetime reproductive performance of domestic animals such as, but not limited to, cows, sheep and pigs.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods.

Assays for activin/inhibin activity include, without limitation, those described in: Vale et al., Endocrinology 91:562-572, 1972; Ling et al., Nature 321:779-782, 1986; Vale et al., Nature 321:776-779, 1986; Mason et al., Nature 318:659-663, 1985; Forage et al., Proc. Natl. Acad. Sci. USA 83:3091-3095, 1986.

4.10.9 CHEMOTACTIC/CHEMOKINETIC ACTIVITY

A polypeptide of the present invention may be involved in chemotactic or chemokinetic activity for mammalian cells, including, for example, monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Chemotactic and chemokinetic receptor activation can be used to mobilize or attract a desired cell population to a desired site of action. Chemotactic or chemokinetic compositions (e.g. proteins, antibodies, binding partners, or modulators of the invention) provide particular advantages in treatment of wounds and other trauma to tissues, as well as in treatment of localized infections. For example, attraction of lymphocytes, monocytes or neutrophils to tumors or sites of infection may result in improved immune responses against the tumor or infecting agent.

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A protein or peptide has chemotactic activity for a particular cell population if it can stimulate, directly or indirectly, the directed orientation or movement of such cell population. Preferably, the protein or peptide has the ability to directly stimulate directed movement of cells. Whether a particular protein has chemotactic activity for a population of cells can be readily determined by employing such protein or peptide in any known assay for cell chemotaxis.

Therapeutic compositions of the invention can be used in the following:

Assays for chemotactic activity (which will identify proteins that induce or prevent chemotaxis) consist of assays that measure the ability of a protein to induce the migration of cells across a membrane as well as the ability of a protein to induce the adhesion of one cell population to another cell population. Suitable assays for movement and adhesion include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Marguiles, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 6.12, Measurement of alpha and beta Chemokines 6.12.1-6.12.28; Taub et al. J. Clin. Invest. 95:1370-1376, 1995; Lind et al. APMIS 103:140-146, 1995; Muller et al Eur. J. Immunol. 25:1744-1748; Gruber et al. J. of Immunol. 152:5860-5867, 1994; Johnston et al. J. of Immunol. 153:1762-1768, 1994.

4.10.10 HEMOSTATIC AND THROMBOLYTIC ACTIVITY

A polypeptide of the invention may also be involved in hemostatis or thrombolysis or thrombosis. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Compositions may be useful in treatment of various coagulation disorders (including hereditary disorders, such as hemophilias) or to enhance coagulation and other hemostatic events in treating wounds resulting from trauma, surgery or other causes. A composition of the invention may also be useful for dissolving or inhibiting formation of thromboses and for treatment and prevention of conditions resulting therefrom (such as, for example, infarction of cardiac and central nervous system vessels (e.g., stroke).

Therapeutic compositions of the invention can be used in the following:

Assay for hemostatic and thrombolytic activity include, without limitation, those described in: Linet et al., J. Clin. Pharmacol. 26:131-140, 1986; Burdick et al., Thrombosis Res. 45:413-419, 1987; Humphrey et al., Fibrinolysis 5:71-79 (1991); Schaub, Prostaglandins 35:467-474, 1988.

4.10.11 CANCER DIAGNOSIS AND THERAPY

Polypeptides of the invention may be involved in cancer cell generation, proliferation or metastasis. Detection of the presence or amount of polynucleotides or polypeptides of the

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invention may be useful for the diagnosis and/or prognosis of one or more types of cancer. For example, the presence or increased expression of a polynucleotide/polypeptide of the invention may indicate a hereditary risk of cancer, a precancerous condition, or an ongoing malignancy. Conversely, a defect in the gene or absence of the polypeptide may be associated with a cancer condition. Identification of single nucleotide polymorphisms associated with cancer or a predisposition to cancer may also be useful for diagnosis or prognosis.

Cancer treatments promote tumor regression by inhibiting tumor cell proliferation, inhibiting angiogenesis (growth of new blood vessels that is necessary to support tumor growth) and/or prohibiting metastasis by reducing tumor cell motility or invasiveness. Therapeutic compositions of the invention may be effective in adult and pediatric oncology including in solid phase tumors/malignancies, locally advanced tumors, human soft tissue sarcomas, metastatic cancer, including lymphatic metastases, blood cell malignancies including multiple myeloma, acute and chronic leukemias, and lymphomas, head and neck cancers including mouth cancer, larynx cancer and thyroid cancer, lung cancers including small cell carcinoma and non-small cell cancers, breast cancers including small cell carcinoma and ductal carcinoma, gastrointestinal cancers including esophageal cancer, stomach cancer, colon cancer, colorectal cancer and polyps associated with colorectal neoplasia, pancreatic cancers, liver cancer, urologic cancers including bladder cancer and prostate cancer, malignancies of the female genital tract including ovarian carcinoma, uterine (including endometrial) cancers, and solid tumor in the ovarian follicle, kidney cancers including renal cell carcinoma, brain cancers including intrinsic brain tumors, neuroblastoma, astrocytic brain tumors, gliomas, metastatic tumor cell invasion in the central nervous system, bone cancers including osteomas, skin cancers including malignant melanoma, tumor progression of human skin keratinocytes, squamous cell carcinoma, basal cell carcinoma, hemangiopericytoma and Karposi's sarcoma.

Polypeptides, polynucleotides, or modulators of polypeptides of the invention (including inhibitors and stimulators of the biological activity of the polypeptide of the invention) may be administered to treat cancer. Therapeutic compositions can be administered in therapeutically effective dosages alone or in combination with adjuvant cancer therapy such as surgery, chemotherapy, radiotherapy, thermotherapy, and laser therapy, and may provide a beneficial effect, *e.g.* reducing tumor size, slowing rate of tumor growth, inhibiting metastasis, or otherwise improving overall clinical condition, without necessarily eradicating the cancer.

The composition can also be administered in therapeutically effective amounts as a portion of an anti-cancer cocktail. An anti-cancer cocktail is a mixture of the polypeptide or modulator of the invention with one or more anti-cancer drugs in addition to a pharmaceutically acceptable carrier for delivery. The use of anti-cancer cocktails as a cancer treatment is routine.

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Anti-cancer drugs that are well known in the art and can be used as a treatment in combination with the polypeptide or modulator of the invention include: Actinomycin D, Aminoglutethimide, Asparaginase, Bleomycin, Busulfan, Carboplatin, Carmustine, Chlorambucil, Cisplatin (cisDDP), Cyclophosphamide, Cytarabine HCl (Cytosine arabinoside), Dacarbazine, Dactinomycin,
Daunorubicin HCl, Doxorubicin HCl, Estramustine phosphate sodium, Etoposide (V16-213), Floxuridine, 5-Fluorouracil (5-Fu), Flutamide, Hydroxyurea (hydroxycarbamide), Ifosfamide, Interferon Alpha-2a, Interferon Alpha-2b, Leuprolide acetate (LHRH-releasing factor analog), Lomustine, Mechlorethamine HCl (nitrogen mustard), Melphalan, Mercaptopurine, Mesna, Methotrexate (MTX), Mitomycin, Mitoxantrone HCl, Octreotide, Plicamycin, Procarbazine HCl, Streptozocin, Tamoxifen citrate, Thioguanine, Thiotepa, Vinblastine sulfate, Vincristine sulfate, Amsacrine, Azacitidine, Hexamethylmelamine, Interleukin-2, Mitoguazone, Pentostatin, Semustine, Teniposide, and Vindesine sulfate.

In addition, therapeutic compositions of the invention may be used for prophylactic treatment of cancer. There are hereditary conditions and/or environmental situations (e.g. exposure to carcinogens) known in the art that predispose an individual to developing cancers. Under these circumstances, it may be beneficial to treat these individuals with therapeutically effective doses of the polypeptide of the invention to reduce the risk of developing cancers.

In vitro models can be used to determine the effective doses of the polypeptide of the invention as a potential cancer treatment. These *in vitro* models include proliferation assays of cultured tumor cells, growth of cultured tumor cells in soft agar (see Freshney, (1987) Culture of Animal Cells: A Manual of Basic Technique, Wily-Liss, New York, NY Ch 18 and Ch 21), tumor systems in nude mice as described in Giovanella et al., J. Natl. Can. Inst., 52: 921-30 (1974), mobility and invasive potential of tumor cells in Boyden Chamber assays as described in Pilkington et al., Anticancer Res., 17: 4107-9 (1997), and angiogenesis assays such as induction of vascularization of the chick chorioallantoic membrane or induction of vascular endothelial cell migration as described in Ribatta et al., Intl. J. Dev. Biol., 40: 1189-97 (1999) and Li et al., Clin. Exp. Metastasis, 17:423-9 (1999), respectively. Suitable tumor cells lines are available, *e.g.* from American Type Tissue Culture Collection catalogs.

4.10.12 RECEPTOR/LIGAND ACTIVITY

A polypeptide of the present invention may also demonstrate activity as receptor, receptor ligand or inhibitor or agonist of receptor/ligand interactions. A polynucleotide of the invention can encode a polypeptide exhibiting such characteristics. Examples of such receptors and ligands include, without limitation, cytokine receptors and their ligands, receptor kinases and their ligands, receptor phosphatases and their ligands, receptors involved in cell-cell interactions

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and their ligands (including without limitation, cellular adhesion molecules (such as selectins, integrins and their ligands) and receptor/ligand pairs involved in antigen presentation, antigen recognition and development of cellular and humoral immune responses. Receptors and ligands are also useful for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. A protein of the present invention (including, without limitation, fragments of receptors and ligands) may themselves be useful as inhibitors of receptor/ligand interactions.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods:

Suitable assays for receptor-ligand activity include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley- Interscience (Chapter 7.28, Measurement of Cellular Adhesion under static conditions 7.28.1-7.28.22), Takai et al., Proc. Natl. Acad. Sci. USA 84:6864-6868, 1987; Bierer et al., J. Exp. Med. 168:1145-1156, 1988; Rosenstein et al., J. Exp. Med. 169:149-160 1989; Stoltenborg et al., J. Immunol. Methods 175:59-68, 1994; Stitt et al., Cell 80:661-670, 1995.

By way of example, the polypeptides of the invention may be used as a receptor for a ligand(s) thereby transmitting the biological activity of that ligand(s). Ligands may be identified through binding assays, affinity chromatography, dihybrid screening assays, BIAcore assays, gel overlay assays, or other methods known in the art.

Studies characterizing drugs or proteins as agonist or antagonist or partial agonists or a partial antagonist require the use of other proteins as competing ligands. The polypeptides of the present invention or ligand(s) thereof may be labeled by being coupled to radioisotopes, colorimetric molecules or a toxin molecules by conventional methods. ("Guide to Protein Purification" Murray P. Deutscher (ed) Methods in Enzymology Vol. 182 (1990) Academic Press, Inc. San Diego). Examples of radioisotopes include, but are not limited to, tritium and carbon-14. Examples of colorimetric molecules include, but are not limited to, fluorescent molecules such as fluorescamine, or rhodamine or other colorimetric molecules. Examples of toxins include, but are not limited, to ricin.

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4.10.13 DRUG SCREENING

This invention is particularly useful for screening chemical compounds by using the novel polypeptides or binding fragments thereof in any of a variety of drug screening techniques. The polypeptides or fragments employed in such a test may either be free in solution, affixed to a solid support, borne on a cell surface or located intracellularly. One method of drug screening

utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or a fragment thereof. Drugs are screened against such transformed cells in competitive binding assays. Such cells, either in viable or fixed form, can be used for standard binding assays. One may measure, for example, the formation of complexes between polypeptides of the invention or fragments and the agent being tested or examine the diminution in complex formation between the novel polypeptides and an appropriate cell line, which are well known in the art.

Sources for test compounds that may be screened for ability to bind to or modulate (i.e., increase or decrease) the activity of polypeptides of the invention include (1) inorganic and organic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of either random or mimetic peptides, oligonucleotides or organic molecules.

Chemical libraries may be readily synthesized or purchased from a number of commercial sources, and may include structural analogs of known compounds or compounds that are identified as "hits" or "leads" via natural product screening.

The sources of natural product libraries are microorganisms (including bacteria and fungi), animals, plants or other vegetation, or marine organisms, and libraries of mixtures for screening may be created by: (1) fermentation and extraction of broths from soil, plant or marine microorganisms or (2) extraction of the organisms themselves. Natural product libraries include polyketides, non-ribosomal peptides, and (non-naturally occurring) variants thereof. For a review, see Science 282:63-68 (1998).

Combinatorial libraries are composed of large numbers of peptides, oligonucleotides or organic compounds and can be readily prepared by traditional automated synthesis methods, PCR, cloning or proprietary synthetic methods. Of particular interest are peptide and oligonucleotide combinatorial libraries. Still other libraries of interest include peptide, protein, peptidomimetic, multiparallel synthetic collection, recombinatorial, and polypeptide libraries. For a review of combinatorial chemistry and libraries created therefrom, see Myers, Curr. Opin. Biotechnol. 8:701-707 (1997). For reviews and examples of peptidomimetic libraries, see Al-Obeidi et al., Mol. Biotechnol, 9(3):205-23 (1998); Hruby et al., Curr Opin Chem Biol, 1(1):114-19 (1997); Dorner et al., Bioorg Med Chem, 4(5):709-15 (1996) (alkylated dipeptides).

Identification of modulators through use of the various libraries described herein permits modification of the candidate "hit" (or "lead") to optimize the capacity of the "hit" to bind a polypeptide of the invention. The molecules identified in the binding assay are then tested for antagonist or agonist activity in in vivo tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

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The binding molecules thus identified may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells such as radioisotopes. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for a polypeptide of the invention. Alternatively, the binding molecules may be complexed with imaging agents for targeting and imaging purposes.

4.10.14 ASSAY FOR RECEPTOR ACTIVITY

The invention also provides methods to detect specific binding of a polypeptide e.g. a ligand or a receptor. The art provides numerous assays particularly useful for identifying previously unknown binding partners for receptor polypeptides of the invention. For example, expression cloning using mammalian or bacterial cells, or dihybrid screening assays can be used to identify polynucleotides encoding binding partners. As another example, affinity chromatography with the appropriate immobilized polypeptide of the invention can be used to isolate polypeptides that recognize and bind polypeptides of the invention. There are a number of different libraries used for the identification of compounds, and in particular small molecules, that modulate (i.e., increase or decrease) biological activity of a polypeptide of the invention. Ligands for receptor polypeptides of the invention can also be identified by adding exogenous ligands, or cocktails of ligands to two cells populations that are genetically identical except for the expression of the receptor of the invention: one cell population expresses the receptor of the invention whereas the other does not. The response of the two cell populations to the addition of ligands(s) are then compared. Alternatively, an expression library can be co-expressed with the polypeptide of the invention in cells and assayed for an autocrine response to identify potential ligand(s). As still another example, BIAcore assays, gel overlay assays, or other methods known in the art can be used to identify binding partner polypeptides, including, (1) organic and inorganic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of random peptides, oligonucleotides or organic molecules.

The role of downstream intracellular signaling molecules in the signaling cascade of the polypeptide of the invention can be determined. For example, a chimeric protein in which the cytoplasmic domain of the polypeptide of the invention is fused to the extracellular portion of a protein, whose ligand has been identified, is produced in a host cell. The cell is then incubated with the ligand specific for the extracellular portion of the chimeric protein, thereby activating the chimeric receptor. Known downstream proteins involved in intracellular signaling can then be assayed for expected modifications *i.e.* phosphorylation. Other methods known to those in the art can also be used to identify signaling molecules involved in receptor activity.

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4.10.15 ANTI-INFLAMMATORY ACTIVITY

Compositions of the present invention may also exhibit anti-inflammatory activity. The anti-inflammatory activity may be achieved by providing a stimulus to cells involved in the inflammatory response, by inhibiting or promoting cell-cell interactions (such as, for example, cell adhesion), by inhibiting or promoting chemotaxis of cells involved in the inflammatory process, inhibiting or promoting cell extravasation, or by stimulating or suppressing production of other factors which more directly inhibit or promote an inflammatory response. Compositions with such activities can be used to treat inflammatory conditions including chronic or acute conditions), including without limitation intimation associated with infection (such as septic shock, sepsis or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine-induced lung injury, inflammatory bowel disease, Crohn's disease or resulting from over production of cytokines such as TNF or IL-1. Compositions of the invention may also be useful to treat anaphylaxis and hypersensitivity to an antigenic substance or material. Compositions of this invention may be utilized to prevent or treat conditions such as, but not limited to, sepsis, acute pancreatitis, endotoxin shock, cytokine induced shock, rheumatoid arthritis, chronic inflammatory arthritis, pancreatic cell damage from diabetes mellitus type 1, graft versus host disease, inflammatory bowel disease, inflamation associated with pulmonary disease, other autoimmune disease or inflammatory disease, an antiproliferative agent such as for acute or chronic mylegenous leukemia or in the prevention of premature labor secondary to intrauterine infections.

4.10.16 LEUKEMIAS

Leukemias and related disorders may be treated or prevented by administration of a

therapeutic that promotes or inhibits function of the polynucleotides and/or polypeptides of the invention. Such leukemias and related disorders include but are not limited to acute leukemia, acute lymphocytic leukemia, acute myelocytic leukemia, myeloblastic, promyelocytic, myelomonocytic, monocytic, erythroleukemia, chronic leukemia, chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia (for a review of such disorders, see

Fishman et al., 1985, Medicine, 2d Ed., J.B. Lippincott Co., Philadelphia).

4.10.17 NERVOUS SYSTEM DISORDERS

Nervous system disorders, involving cell types which can be tested for efficacy of intervention with compounds that modulate the activity of the polynucleotides and/or polypeptides of the invention, and which can be treated upon thus observing an indication of

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therapeutic utility, include but are not limited to nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and non-human mammalian patients) according to the invention include but are not limited to the following lesions of either the central (including spinal cord, brain) or peripheral nervous systems:

- (i) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries;
- (ii) ischemic lesions, in which a lack of oxygen in a portion of the nervous system results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord infarction or ischemia;
- (iii) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, syphilis;
- (iv) degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis;
- (v) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration;
- (vi) neurological lesions associated with systemic diseases including but not limited to diabetes (diabetic neuropathy, Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis;
- (vii) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and
 - (viii) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including but not limited to multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

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Therapeutics which are useful according to the invention for treatment of a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way of limitation, therapeutics which elicit any of the following effects may be useful according to the invention:

- (i) increased survival time of neurons in culture;
- (ii) increased sprouting of neurons in culture or in vivo;
- (iii) increased production of a neuron-associated molecule in culture or *in vivo*, *e.g.*, choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or
 - (iv) decreased symptoms of neuron dysfunction in vivo.

Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may be measured by the method set forth in Arakawa et al. (1990, J. Neurosci. 10:3507-3515); increased sprouting of neurons may be detected by methods set forth in Pestronk et al. (1980, Exp. Neurol. 70:65-82) or Brown et al. (1981, Ann. Rev. Neurosci. 4:17-42); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay, etc., depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

In specific embodiments, motor neuron disorders that may be treated according to the invention include but are not limited to disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including but not limited to progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome), poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

4.10.18 OTHER ACTIVITIES

A polypeptide of the invention may also exhibit one or more of the following additional activities or effects: inhibiting the growth, infection or function of, or killing, infectious agents, including, without limitation, bacteria, viruses, fungi and other parasites; effecting (suppressing or enhancing) bodily characteristics, including, without limitation, height, weight, hair color, eye color, skin, fat to lean ratio or other tissue pigmentation, or organ or body part size or shape (such as, for example, breast augmentation or diminution, change in bone form or shape);

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effecting biorhythms or circadian cycles or rhythms; effecting the fertility of male or female subjects; effecting the metabolism, catabolism, anabolism, processing, utilization, storage or elimination of dietary fat, lipid, protein, carbohydrate, vitamins, minerals, co-factors or other nutritional factors or component(s); effecting behavioral characteristics, including, without limitation, appetite, libido, stress, cognition (including cognitive disorders), depression (including depressive disorders) and violent behaviors; providing analgesic effects or other pain reducing effects; promoting differentiation and growth of embryonic stem cells in lineages other than hematopoietic lineages; hormonal or endocrine activity; in the case of enzymes, correcting deficiencies of the enzyme and treating deficiency-related diseases; treatment of hyperproliferative disorders (such as, for example, psoriasis); immunoglobulin-like activity (such as, for example, the ability to bind antigens or complement); and the ability to act as an antigen in a vaccine composition to raise an immune response against such protein or another material or entity which is cross-reactive with such protein.

4.10.19 IDENTIFICATION OF POLYMORPHISMS

The demonstration of polymorphisms makes possible the identification of such polymorphisms in human subjects and the pharmacogenetic use of this information for diagnosis and treatment. Such polymorphisms may be associated with, *e.g.*, differential predisposition or susceptibility to various disease states (such as disorders involving inflammation or immune response) or a differential response to drug administration, and this genetic information can be used to tailor preventive or therapeutic treatment appropriately. For example, the existence of a polymorphism associated with a predisposition to inflammation or autoimmune disease makes possible the diagnosis of this condition in humans by identifying the presence of the polymorphism.

Polymorphisms can be identified in a variety of ways known in the art which all generally involve obtaining a sample from a patient, analyzing DNA from the sample, optionally involving isolation or amplification of the DNA, and identifying the presence of the polymorphism in the DNA. For example, PCR may be used to amplify an appropriate fragment of genomic DNA which may then be sequenced. Alternatively, the DNA may be subjected to allele-specific oligonucleotide hybridization (in which appropriate oligonucleotides are hybridized to the DNA under conditions permitting detection of a single base mismatch) or to a single nucleotide extension assay (in which an oligonucleotide that hybridizes immediately adjacent to the position of the polymorphism is extended with one or more labeled nucleotides). In addition, traditional restriction fragment length polymorphism analysis (using restriction enzymes that provide differential digestion of the genomic DNA depending on the presence or

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absence of the polymorphism) may be performed. Arrays with nucleotide sequences of the present invention can be used to detect polymorphisms. The array can comprise modified nucleotide sequences of the present invention in order to detect the nucleotide sequences of the present invention. In the alternative, any one of the nucleotide sequences of the present invention can be placed on the array to detect changes from those sequences.

Alternatively a polymorphism resulting in a change in the amino acid sequence could also be detected by detecting a corresponding change in amino acid sequence of the protein, e.g., by an antibody specific to the variant sequence.

4.10.20 ARTHRITIS AND INFLAMMATION

The immunosuppressive effects of the compositions of the invention against rheumatoid arthritis is determined in an experimental animal model system. The experimental model system is adjuvant induced arthritis in rats, and the protocol is described by J. Holoshitz, et at., 1983, Science, 219:56, or by B. Waksman et al., 1963, Int. Arch. Allergy Appl. Immunol., 23:129. Induction of the disease can be caused by a single injection, generally intradermally, of a suspension of killed Mycobacterium tuberculosis in complete Freund's adjuvant (CFA). The route of injection can vary, but rats may be injected at the base of the tail with an adjuvant mixture. The polypeptide is administered in phosphate buffered solution (PBS) at a dose of about 1-5 mg/kg. The control consists of administering PBS only.

The procedure for testing the effects of the test compound would consist of intradermally injecting killed Mycobacterium tuberculosis in CFA followed by immediately administering the test compound and subsequent treatment every other day until day 24. At 14, 15, 18, 20, 22, and 24 days after injection of Mycobacterium CFA, an overall arthritis score may be obtained as described by J. Holoskitz above. An analysis of the data would reveal that the test compound would have a dramatic affect on the swelling of the joints as measured by a decrease of the arthritis score.

4.11 THERAPEUTIC METHODS

The compositions (including polypeptide fragments, analogs, variants and antibodies or other binding partners or modulators including antisense polynucleotides) of the invention have numerous applications in a variety of therapeutic methods. Examples of therapeutic applications include, but are not limited to, those exemplified herein.

4.11.1 EXAMPLE

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One embodiment of the invention is the administration of an effective amount of the polypeptides or other composition of the invention to individuals affected by a disease or disorder that can be modulated by regulating the peptides of the invention. While the mode of administration is not particularly important, parenteral administration is preferred. An exemplary mode of administration is to deliver an intravenous bolus. The dosage of the polypeptides or other composition of the invention will normally be determined by the prescribing physician. It is to be expected that the dosage will vary according to the age, weight, condition and response of the individual patient. Typically, the amount of polypeptide administered per dose will be in the range of about 0.01µg/kg to 100 mg/kg of body weight, with the preferred dose being about 0.1µg/kg to 10 mg/kg of patient body weight. For parenteral administration, polypeptides of the invention will be formulated in an injectable form combined with a pharmaceutically acceptable parenteral vehicle. Such vehicles are well known in the art and examples include water, saline, Ringer's solution, dextrose solution, and solutions consisting of small amounts of the human serum albumin. The vehicle may contain minor amounts of additives that maintain the isotonicity and stability of the polypeptide or other active ingredient. The preparation of such solutions is within the skill of the art.

4.12 PHARMACEUTICAL FORMULATIONS AND ROUTES OF ADMINISTRATION

A protein or other composition of the present invention (from whatever source derived, including without limitation from recombinant and non-recombinant sources and including antibodies and other binding partners of the polypeptides of the invention) may be administered to a patient in need, by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s) at doses to treat or ameliorate a variety of disorders. Such a composition may optionally contain (in addition to protein or other active ingredient and a carrier) diluents. fillers, salts, buffers, stabilizers, solubilizers, and other materials well known in the art. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredient(s). The characteristics of the carrier will depend on the route of administration. The pharmaceutical composition of the invention may also contain cytokines, lymphokines, or other hematopoietic factors such as M-CSF, GM-CSF, TNF, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IFN, TNF0, TNF1, TNF2, G-CSF, Meg-CSF, thrombopoietin, stem cell factor, and erythropoietin. In further compositions, proteins of the invention may be combined with other agents beneficial to the treatment of the disease or disorder in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet-derived growth

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factor (PDGF), transforming growth factors (TGF- α and TGF- β), insulin-like growth factor (IGF), as well as cytokines described herein.

The pharmaceutical composition may further contain other agents which either enhance the activity of the protein or other active ingredient or complement its activity or use in treatment. Such additional factors and/or agents may be included in the pharmaceutical composition to produce a synergistic effect with protein or other active ingredient of the invention, or to minimize side effects. Conversely, protein or other active ingredient of the present invention may be included in formulations of the particular clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent to minimize side effects of the clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent (such as IL-1Ra, IL-1 Hy1, IL-1 Hy2, anti-TNF, corticosteroids, immunosuppressive agents). A protein of the present invention may be active in multimers (e.g., heterodimers or homodimers) or complexes with itself or other proteins. As a result, pharmaceutical compositions of the invention may comprise a protein of the invention in such multimeric or complexed form.

As an alternative to being included in a pharmaceutical composition of the invention including a first protein, a second protein or a therapeutic agent may be concurrently administered with the first protein (e.g., at the same time, or at differing times provided that therapeutic concentrations of the combination of agents is achieved at the treatment site). Techniques for formulation and administration of the compounds of the instant application may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, PA, latest edition. A therapeutically effective dose further refers to that amount of the compound sufficient to result in amelioration of symptoms, e.g., treatment, healing, prevention or amelioration of the relevant medical condition, or an increase in rate of treatment, healing, prevention or amelioration of such conditions. When applied to an individual active ingredient, administered alone, a therapeutically effective dose refers to that ingredient alone. When applied to a combination, a therapeutically effective dose refers to combined amounts of the active ingredients that result in the therapeutic effect, whether administered in combination, serially or simultaneously.

In practicing the method of treatment or use of the present invention, a therapeutically effective amount of protein or other active ingredient of the present invention is administered to a mammal having a condition to be treated. Protein or other active ingredient of the present invention may be administered in accordance with the method of the invention either alone or in combination with other therapies such as treatments employing cytokines, lymphokines or other hematopoietic factors. When co- administered with one or more cytokines, lymphokines or other

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hematopoietic factors, protein or other active ingredient of the present invention may be administered either simultaneously with the cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors, or sequentially. If administered sequentially, the attending physician will decide on the appropriate sequence of administering protein or other active ingredient of the present invention in combination with cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors.

4.12.1 ROUTES OF ADMINISTRATION

Suitable routes of administration may, for example, include oral, rectal, transmucosal, or intestinal administration; parenteral delivery, including intramuscular, subcutaneous, intramedullary injections, as well as intrathecal, direct intraventricular, intravenous, intraperitoneal, intranasal, or intraocular injections. Administration of protein or other active ingredient of the present invention used in the pharmaceutical composition or to practice the method of the present invention can be carried out in a variety of conventional ways, such as oral ingestion, inhalation, topical application or cutaneous, subcutaneous, intraperitoneal, parenteral or intravenous injection. Intravenous administration to the patient is preferred.

Alternately, one may administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into a arthritic joints or in fibrotic tissue, often in a depot or sustained release formulation. In order to prevent the scarring process frequently occurring as complication of glaucoma surgery, the compounds may be administered topically, for example, as eye drops. Furthermore, one may administer the drug in a targeted drug delivery system, for example, in a liposome coated with a specific antibody, targeting, for example, arthritic or fibrotic tissue. The liposomes will be targeted to and taken up selectively by the afflicted tissue.

The polypeptides of the invention are administered by any route that delivers an effective dosage to the desired site of action. The determination of a suitable route of administration and an effective dosage for a particular indication is within the level of skill in the art. Preferably for wound treatment, one administers the therapeutic compound directly to the site. Suitable dosage ranges for the polypeptides of the invention can be extrapolated from these dosages or from similar studies in appropriate animal models. Dosages can then be adjusted as necessary by the clinician to provide maximal therapeutic benefit.

4.12.2 COMPOSITIONS/FORMULATIONS

Pharmaceutical compositions for use in accordance with the present invention thus may be formulated in a conventional manner using one or more physiologically acceptable carriers

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comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. These pharmaceutical compositions may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Proper formulation is dependent upon the route of administration chosen. When a therapeutically effective amount of protein or other active ingredient of the present invention is administered orally, protein or other active ingredient of the present invention will be in the form of a tablet, capsule, powder, solution or elixir. When administered in tablet form, the pharmaceutical composition of the invention may additionally contain a solid carrier such as a gelatin or an adjuvant. The tablet, capsule, and powder contain from about 5 to 95% protein or other active ingredient of the present invention, and preferably from about 25 to 90% protein or other active ingredient of the present invention. When administered in liquid form, a liquid carrier such as water, petroleum, oils of animal or plant origin such as peanut oil, mineral oil, soybean oil, or sesame oil, or synthetic oils may be added. The liquid form of the pharmaceutical composition may further contain physiological saline solution, dextrose or other saccharide solution, or glycols such as ethylene glycol, propylene glycol or polyethylene glycol. When administered in liquid form, the pharmaceutical composition contains from about 0.5 to 90% by weight of protein or other active ingredient of the present invention, and preferably from about 1 to 50% protein or other active ingredient of the present invention.

When a therapeutically effective amount of protein or other active ingredient of the present invention is administered by intravenous, cutaneous or subcutaneous injection, protein or other active ingredient of the present invention will be in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable protein or other active ingredient solutions, having due regard to pH, isotonicity, stability, and the like, is within the skill in the art. A preferred pharmaceutical composition for intravenous, cutaneous, or subcutaneous injection should contain, in addition to protein or other active ingredient of the present invention, an isotonic vehicle such as Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection, or other vehicle as known in the art. The pharmaceutical composition of the present invention may also contain stabilizers, preservatives, buffers, antioxidants, or other additives known to those of skill in the art. For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

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For oral administration, the compounds can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained from a solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate. Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration. For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebuliser, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch. The compounds may be formulated for parenteral

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administration by injection, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides. In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

A pharmaceutical carrier for the hydrophobic compounds of the invention is a co-solvent system comprising benzyl alcohol, a nonpolar surfactant, a water-miscible organic polymer, and an aqueous phase. The co-solvent system may be the VPD co-solvent system. VPD is a solution of 3% w/v benzyl alcohol, 8% w/v of the nonpolar surfactant polysorbate 80, and 65% w/v polyethylene glycol 300, made up to volume in absolute ethanol. The VPD co-solvent system (VPD:5W) consists of VPD diluted 1:1 with a 5% dextrose in water solution. This co-solvent system dissolves hydrophobic compounds well, and itself produces low toxicity upon systemic administration. Naturally, the proportions of a co-solvent system may be varied considerably without destroying its solubility and toxicity characteristics. Furthermore, the identity of the co-solvent components may be varied: for example, other low-toxicity nonpolar surfactants may be used instead of polysorbate 80; the fraction size of polyethylene glycol may be varied; other biocompatible polymers may replace polyethylene glycol, e.g. polyvinyl pyrrolidone; and other

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sugars or polysaccharides may substitute for dextrose. Alternatively, other delivery systems for hydrophobic pharmaceutical compounds may be employed. Liposomes and emulsions are well known examples of delivery vehicles or carriers for hydrophobic drugs. Certain organic solvents such as dimethylsulfoxide also may be employed, although usually at the cost of greater toxicity. Additionally, the compounds may be delivered using a sustained-release system, such as semipermeable matrices of solid hydrophobic polymers containing the therapeutic agent. Various types of sustained-release materials have been established and are well known by those skilled in the art. Sustained-release capsules may, depending on their chemical nature, release the compounds for a few weeks up to over 100 days. Depending on the chemical nature and the biological stability of the therapeutic reagent, additional strategies for protein or other active ingredient stabilization may be employed.

The pharmaceutical compositions also may comprise suitable solid or gel phase carriers or excipients. Examples of such carriers or excipients include but are not limited to calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols. Many of the active ingredients of the invention may be provided as salts with pharmaceutically compatible counter ions. Such pharmaceutically acceptable base addition salts are those salts which retain the biological effectiveness and properties of the free acids and which are obtained by reaction with inorganic or organic bases such as sodium hydroxide, magnesium hydroxide, ammonia, trialkylamine, dialkylamine, monoalkylamine, dibasic amino acids, sodium acetate, potassium benzoate, triethanol amine and the like.

The pharmaceutical composition of the invention may be in the form of a complex of the protein(s) or other active ingredient(s) of present invention along with protein or peptide antigens. The protein and/or peptide antigen will deliver a stimulatory signal to both B and T lymphocytes. B lymphocytes will respond to antigen through their surface immunoglobulin receptor. T lymphocytes will respond to antigen through the T cell receptor (TCR) following presentation of the antigen by MHC proteins. MHC and structurally related proteins including those encoded by class I and class II MHC genes on host cells will serve to present the peptide antigen(s) to T lymphocytes. The antigen components could also be supplied as purified MHC-peptide complexes alone or with co-stimulatory molecules that can directly signal T cells. Alternatively antibodies able to bind surface immunoglobulin and other molecules on B cells as well as antibodies able to bind the TCR and other molecules on T cells can be combined with the pharmaceutical composition of the invention.

The pharmaceutical composition of the invention may be in the form of a liposome in which protein of the present invention is combined, in addition to other pharmaceutically

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acceptable carriers, with amphipathic agents such as lipids which exist in aggregated form as micelles, insoluble monolayers, liquid crystals, or lamellar layers in aqueous solution. Suitable lipids for liposomal formulation include, without limitation, monoglycerides, diglycerides, sulfatides, lysolecithins, phospholipids, saponin, bile acids, and the like. Preparation of such liposomal formulations is within the level of skill in the art, as disclosed, for example, in U.S. Patent Nos. 4,235,871; 4,501,728; 4,837,028; and 4,737,323, all of which are incorporated herein by reference.

The amount of protein or other active ingredient of the present invention in the pharmaceutical composition of the present invention will depend upon the nature and severity of the condition being treated, and on the nature of prior treatments which the patient has undergone. Ultimately, the attending physician will decide the amount of protein or other active ingredient of the present invention with which to treat each individual patient. Initially, the attending physician will administer low doses of protein or other active ingredient of the present invention and observe the patient's response. Larger doses of protein or other active ingredient of the present invention may be administered until the optimal therapeutic effect is obtained for the patient, and at that point the dosage is not increased further. It is contemplated that the various pharmaceutical compositions used to practice the method of the present invention should contain about 0.01 µg to about 100 mg (preferably about 0.1 µg to about 10 mg, more preferably about 0.1 µg to about 1 mg) of protein or other active ingredient of the present invention per kg body weight. For compositions of the present invention which are useful for bone, cartilage, tendon or ligament regeneration, the therapeutic method includes administering the composition topically, systematically, or locally as an implant or device. When administered, the therapeutic composition for use in this invention is, of course, in a pyrogen-free, physiologically acceptable form. Further, the composition may desirably be encapsulated or injected in a viscous form for delivery to the site of bone, cartilage or tissue damage. Topical administration may be suitable for wound healing and tissue repair. Therapeutically useful agents other than a protein or other active ingredient of the invention which may also optionally be included in the composition as described above, may alternatively or additionally, be administered simultaneously or sequentially with the composition in the methods of the invention. Preferably for bone and/or cartilage formation, the composition would include a matrix capable of delivering the protein-containing or other active ingredient-containing composition to the site of bone and/or cartilage damage, providing a structure for the developing bone and cartilage and optimally capable of being resorbed into the body. Such matrices may be formed of materials presently in use for other implanted medical applications.

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The choice of matrix material is based on biocompatibility, biodegradability, mechanical. properties, cosmetic appearance and interface properties. The particular application of the compositions will define the appropriate formulation. Potential matrices for the compositions may be biodegradable and chemically defined calcium sulfate, tricalcium phosphate, hydroxyapatite, polylactic acid, polyglycolic acid and polyanhydrides. Other potential materials are biodegradable and biologically well-defined, such as bone or dermal collagen. Further matrices are comprised of pure proteins or extracellular matrix components. Other potential matrices are nonbiodegradable and chemically defined, such as sintered hydroxyapatite, bioglass. aluminates, or other ceramics. Matrices may be comprised of combinations of any of the above mentioned types of material, such as polylactic acid and hydroxyapatite or collagen and tricalcium phosphate. The bioceramics may be altered in composition, such as in calcium-aluminate-phosphate and processing to alter pore size, particle size, particle shape, and biodegradability. Presently preferred is a 50:50 (mole weight) copolymer of lactic acid and glycolic acid in the form of porous particles having diameters ranging from 150 to 800 microns. In some applications, it will be useful to utilize a sequestering agent, such as carboxymethyl cellulose or autologous blood clot, to prevent the protein compositions from disassociating from the matrix.

A preferred family of sequestering agents is cellulosic materials such as alkylcelluloses (including hydroxyalkylcelluloses), including methylcellulose, ethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropyl-methylcellulose, and carboxymethylcellulose, the most preferred being cationic salts of carboxymethylcellulose (CMC). Other preferred sequestering agents include hyaluronic acid, sodium alginate, poly(ethylene glycol), polyoxyethylene oxide, carboxyvinyl polymer and poly(vinyl alcohol). The amount of sequestering agent useful herein is 0.5-20 wt %, preferably 1-10 wt % based on total formulation weight, which represents the amount necessary to prevent desorption of the protein from the polymer matrix and to provide appropriate handling of the composition, yet not so much that the progenitor cells are prevented from infiltrating the matrix, thereby providing the protein the opportunity to assist the osteogenic activity of the progenitor cells. In further compositions, proteins or other active ingredients of the invention may be combined with other agents beneficial to the treatment of the bone and/or cartilage defect, wound, or tissue in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet derived growth factor (PDGF), transforming growth factors (TGF-α and TGF-β), and insulin-like growth factor (IGF).

The therapeutic compositions are also presently valuable for veterinary applications. Particularly domestic animals and thoroughbred horses, in addition to humans, are desired

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patients for such treatment with proteins or other active ingredients of the present invention. The dosage regimen of a protein-containing pharmaceutical composition to be used in tissue regeneration will be determined by the attending physician considering various factors which modify the action of the proteins, *e.g.*, amount of tissue weight desired to be formed, the site of damage, the condition of the damaged tissue, the size of a wound, type of damaged tissue (*e.g.*, bone), the patient's age, sex, and diet, the severity of any infection, time of administration and other clinical factors. The dosage may vary with the type of matrix used in the reconstitution and with inclusion of other proteins in the pharmaceutical composition. For example, the addition of other known growth factors, such as IGF I (insulin like growth factor I), to the final composition, may also effect the dosage. Progress can be monitored by periodic assessment of tissue/bone growth and/or repair, for example, X-rays, histomorphometric determinations and tetracycline labeling.

Polynucleotides of the present invention can also be used for gene therapy. Such polynucleotides can be introduced either in vivo or ex vivo into cells for expression in a mammalian subject. Polynucleotides of the invention may also be administered by other known methods for introduction of nucleic acid into a cell or organism (including, without limitation, in the form of viral vectors or naked DNA). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes.

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4.12.3 EFFECTIVE DOSAGE

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve its intended purpose. More specifically, a therapeutically effective amount means an amount effective to prevent development of or to alleviate the existing symptoms of the subject being treated. Determination of the effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from appropriate in vitro assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that can be used to more accurately determine useful doses in humans. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC₅₀ as determined in cell culture (*i.e.*, the concentration of the test compound which achieves a half-maximal inhibition of the protein's biological activity). Such information can be used to more accurately determine useful doses in humans.

A therapeutically effective dose refers to that amount of the compound that results in amelioration of symptoms or a prolongation of survival in a patient. Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED_{50} (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio between LD₅₀ and ED₅₀. Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. See, e.g., Fingl et al., 1975, in "The Pharmacological Basis of Therapeutics", Ch. 1 p.1. Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain the desired effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from in vitro data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays can be used to determine plasma concentrations.

Dosage intervals can also be determined using MEC value. Compounds should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%. In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

An exemplary dosage regimen for polypeptides or other compositions of the invention will be in the range of about $0.01~\mu g/kg$ to 100~mg/kg of body weight daily, with the preferred dose being about $0.1~\mu g/kg$ to 25~mg/kg of patient body weight daily, varying in adults and children. Dosing may be once daily, or equivalent doses may be delivered at longer or shorter intervals.

The amount of composition administered will, of course, be dependent on the subject being treated, on the subject's age and weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

4.12.4 PACKAGING

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The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. Compositions comprising a compound of the invention formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of an indicated condition.

4.13 ANTIBODIES

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Also included in the invention are antibodies to proteins, or fragments of proteins of the invention. The term "antibody" as used herein refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, *i.e.*, molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, F_{ab} , F_{ab} , and $F_{(ab)2}$ fragments, and an F_{ab} expression library. In general, an antibody molecule obtained from humans relates to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG_1 , IgG_2 , and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain. Reference herein to antibodies includes a reference to all such classes, subclasses and types of human antibody species.

An isolated related protein of the invention may be intended to serve as an antigen, or a portion or fragment thereof, and additionally can be used as an immunogen to generate antibodies that immunospecifically bind the antigen, using standard techniques for polyclonal and monoclonal antibody preparation. The full-length protein can be used or, alternatively, the invention provides antigenic peptide fragments of the antigen for use as immunogens. An antigenic peptide fragment comprises at least 6 amino acid residues of the amino acid sequence of the full length protein, (for example the amino acid sequence shown in SEQ ID NO: 1010), and encompasses an epitope thereof such that an antibody raised against the peptide forms a specific immune complex with the full length protein or with any fragment that contains the epitope. Preferably, the antigenic peptide comprises at least 10 amino acid residues, or at least 15 amino acid residues, or at least 20 amino acid residues. Preferred epitopes encompassed by the antigenic peptide are regions of the protein that are located on its surface; commonly these are hydrophilic regions.

In certain embodiments of the invention, at least one epitope encompassed by the antigenic peptide is a region of -related protein that is located on the surface of the protein, e.g., a hydrophilic region. A hydrophobicity analysis of the human related protein sequence will

indicate which regions of a related protein are particularly hydrophilic and, therefore, are likely to encode surface residues useful for targeting antibody production. As a means for targeting antibody production, hydropathy plots showing regions of hydrophilicity and hydrophobicity may be generated by any method well known in the art, including, for example, the Kyte Doolittle or the Hopp Woods methods, either with or without Fourier transformation. See, *e.g.*, Hopp and Woods, 1981, *Proc. Nat. Acad. Sci. USA* 78: 3824-3828; Kyte and Doolittle 1982, *J. Mol. Biol.* 157: 105-142, each of which is incorporated herein by reference in its entirety. Antibodies that are specific for one or more domains within an antigenic protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

A protein of the invention, or a derivative, fragment, analog, homolog or ortholog thereof, may be utilized as an immunogen in the generation of antibodies that immunospecifically bind these protein components.

Various procedures known within the art may be used for the production of polyclonal or monoclonal antibodies directed against a protein of the invention, or against derivatives, fragments, analogs homologs or orthologs thereof (see, for example, Antibodies: A Laboratory Manual, Harlow E, and Lane D, 1988, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, incorporated herein by reference). Some of these antibodies are discussed below.

5.13.1 Polyclonal Antibodies

For the production of polyclonal antibodies, various suitable host animals (e.g., rabbit, goat, mouse or other mammal) may be immunized by one or more injections with the native protein, a synthetic variant thereof, or a derivative of the foregoing. An appropriate immunogenic preparation can contain, for example, the naturally occurring immunogenic protein, a chemically synthesized polypeptide representing the immunogenic protein, or a recombinantly expressed immunogenic protein. Furthermore, the protein may be conjugated to a second protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. The preparation can further include an adjuvant. Various adjuvants used to increase the immunological response include, but are not limited to, Freund's (complete and incomplete), mineral gels (e.g., aluminum hydroxide), surface active substances (e.g., lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, dinitrophenol, etc.), adjuvants usable in humans such as Bacille Calmette-Guerin and Corynebacterium parvum, or similar immunostimulatory agents. Additional examples of adjuvants which can be employed include MPL-TDM adjuvant (monophosphoryl Lipid A. synthetic trehalose dicorynomycolate).

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The polyclonal antibody molecules directed against the immunogenic protein can be isolated from the mammal (e.g., from the blood) and further purified by well known techniques, such as affinity chromatography using protein A or protein G, which provide primarily the IgG fraction of immune serum. Subsequently, or alternatively, the specific antigen which is the target of the immunoglobulin sought, or an epitope thereof, may be immobilized on a column to purify the immune specific antibody by immunoaffinity chromatography. Purification of immunoglobulins is discussed, for example, by D. Wilkinson (The Scientist, published by The Scientist, Inc., Philadelphia PA, Vol. 14, No. 8 (April 17, 2000), pp. 25-28).

5.13.2 Monoclonal Antibodies

The term "monoclonal antibody" (MAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs thus contain an antigen binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

Monoclonal antibodies can be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes can be immunized in vitro.

The immunizing agent will typically include the protein antigen, a fragment thereof or a fusion protein thereof. Generally, either peripheral blood lymphocytes are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, Monoclonal Antibodies: Principles and Practice, Academic Press, (1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells can be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

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Preferred immortalized cell lines are those that fuse efficiently, support stable high level

expression of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Manassas, Virginia. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, J. Immunol., 133:3001 (1984); Brodeur et al., Monoclonal Antibody Production Techniques and Applications, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the antigen. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, <u>Anal. Biochem.</u>, <u>107</u>:220 (1980). Preferably, antibodies having a high degree of specificity and a high binding affinity for the target antigen are isolated.

After the desired hybridoma cells are identified, the clones can be subcloned by limiting dilution procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells can be grown in vivo as ascites in a mammal.

The monoclonal antibodies secreted by the subclones can be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies can also be made by recombinant DNA methods, such as those described in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA can be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also can be modified, for

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example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (U.S. Patent No. 4,816,567; Morrison, Nature 368, 812-13 (1994)) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

5.13.2 Humanized Antibodies

10 The antibodies directed against the protein antigens of the invention can further comprise humanized antibodies or human antibodies. These antibodies are suitable for administration to humans without engendering an immune response by the human against the administered immunoglobulin. Humanized forms of antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-15 binding subsequences of antibodies) that are principally comprised of the sequence of a human immunoglobulin, and contain minimal sequence derived from a non-human immunoglobulin. Humanization can be performed following the method of Winter and co-workers (Jones et al., Nature, 321:522-525 (1986); Riechmann et al., Nature, 332:323-327 (1988); Verhoeyen et al., Science, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the 20 corresponding sequences of a human antibody. (See also U.S. Patent No. 5,225,539.) In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies can also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable 25 domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., 1986; Riechmann et al., 1988; and Presta, Curr. Op. Struct. Biol., 30 2:593-596 (1992)).

5.13.3 Human Antibodies

Fully human antibodies relate to antibody molecules in which essentially the entire sequences of both the light chain and the heavy chain, including the CDRs, arise from human genes. Such antibodies are termed "human antibodies", or "fully human antibodies" herein.

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Human monoclonal antibodies can be prepared by the trioma technique; the human B-cell hybridoma technique (see Kozbor, et al., 1983 Immunol Today 4: 72) and the EBV hybridoma technique to produce human monoclonal antibodies (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96). Human monoclonal antibodies may be utilized in the practice of the present invention and may be produced by using human hybridomas (see Cote, et al., 1983. Proc Natl Acad Sci USA 80: 2026-2030) or by transforming human B-cells with Epstein Barr Virus in vitro (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96).

In addition, human antibodies can also be produced using additional techniques, including phage display libraries (Hoogenboom and Winter, <u>J. Mol. Biol.</u>, <u>227</u>:381 (1991); Marks et al., <u>J. Mol. Biol.</u>, <u>222</u>:581 (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, *e.g.*, mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, for example, in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in Marks et al. (<u>Bio/Technology 10</u>, 779-783 (1992)); Lonberg et al. (<u>Nature 368</u> 856-859 (1994)); Morrison (<u>Nature 368</u>, 812-13 (1994)); Fishwild et al.(<u>Nature Biotechnology 14</u>, 845-51 (1996)); Neuberger (<u>Nature Biotechnology 14</u>, 826 (1996)); and Lonberg and Huszar (Intern. Rev. Immunol. 13 65-93 (1995)).

Human antibodies may additionally be produced using transgenic nonhuman animals which are modified so as to produce fully human antibodies rather than the animal's endogenous antibodies in response to challenge by an antigen. (See PCT publication WO94/02602). The endogenous genes encoding the heavy and light immunoglobulin chains in the nonhuman host have been incapacitated, and active loci encoding human heavy and light chain immunoglobulins are inserted into the host's genome. The human genes are incorporated, for example, using yeast artificial chromosomes containing the requisite human DNA segments. An animal which provides all the desired modifications is then obtained as progeny by crossbreeding intermediate transgenic animals containing fewer than the full complement of the modifications. The preferred embodiment of such a nonhuman animal is a mouse, and is termed the XenomouseTM as disclosed in PCT publications WO 96/33735 and WO 96/34096. This animal produces B cells which secrete fully human immunoglobulins. The antibodies can be obtained directly from the animal after immunization with an immunogen of interest, as, for example, a preparation of a polyclonal antibody, or alternatively from immortalized B cells derived from the animal, such as hybridomas producing monoclonal antibodies. Additionally, the genes encoding the

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immunoglobulins with human variable regions can be recovered and expressed to obtain the antibodies directly, or can be further modified to obtain analogs of antibodies such as, for example, single chain Fv molecules.

An example of a method of producing a nonhuman host, exemplified as a mouse, lacking expression of an endogenous immunoglobulin heavy chain is disclosed in U.S. Patent No. 5,939,598. It can be obtained by a method including deleting the J segment genes from at least one endogenous heavy chain locus in an embryonic stem cell to prevent rearrangement of the locus and to prevent formation of a transcript of a rearranged immunoglobulin heavy chain locus, the deletion being effected by a targeting vector containing a gene encoding a selectable marker; and producing from the embryonic stem cell a transgenic mouse whose somatic and germ cells contain the gene encoding the selectable marker.

A method for producing an antibody of interest, such as a human antibody, is disclosed in U.S. Patent No. 5,916,771. It includes introducing an expression vector that contains a nucleotide sequence encoding a heavy chain into one mammalian host cell in culture, introducing an expression vector containing a nucleotide sequence encoding a light chain into another mammalian host cell, and fusing the two cells to form a hybrid cell. The hybrid cell expresses an antibody containing the heavy chain and the light chain.

In a further improvement on this procedure, a method for identifying a clinically relevant epitope on an immunogen, and a correlative method for selecting an antibody that binds immunospecifically to the relevant epitope with high affinity, are disclosed in PCT publication WO 99/53049.

5.13.4 Fab Fragments and Single Chain Antibodies

According to the invention, techniques can be adapted for the production of single-chain antibodies specific to an antigenic protein of the invention (see e.g., U.S. Patent No. 4,946,778). In addition, methods can be adapted for the construction of F_{ab} expression libraries (see e.g., Huse, et al., 1989 Science 246: 1275-1281) to allow rapid and effective identification of monoclonal F_{ab} fragments with the desired specificity for a protein or derivatives, fragments, analogs or homologs thereof. Antibody fragments that contain the idiotypes to a protein antigen may be produced by techniques known in the art including, but not limited to: (i) an $F_{(ab')2}$ fragment produced by pepsin digestion of an antibody molecule; (ii) an F_{ab} fragment generated by reducing the disulfide bridges of an $F_{(ab')2}$ fragment; (iii) an F_{ab} fragment generated by the treatment of the antibody molecule with papain and a reducing agent and (iv) F_v fragments.

5.13.5 Bispecific Antibodies

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Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for an antigenic protein of the invention. The second binding target is any other antigen, and advantageously is a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities (Milstein and Cuello, Nature, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker *et al.*, 1991 *EMBO J.*, 10:3655-3659.

Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. For further details of generating bispecific antibodies see, for example, Suresh et al., Methods in Enzymology, 121:210 (1986).

According to another approach described in WO 96/27011, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers which are recovered from recombinant cell culture. The preferred interface comprises at least a part of the CH3 region of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g. tyrosine or tryptophan). Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g. alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers.

Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g. F(ab')₂ bispecific antibodies). Techniques for generating bispecific antibodies from antibody fragments have been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., Science 229:81 (1985) describe a procedure

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wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

Additionally, Fab' fragments can be directly recovered from E. coli and chemically coupled to form bispecific antibodies. Shalaby et al., J. Exp. Med. 175:217-225 (1992) describe the production of a fully humanized bispecific antibody F(ab')₂ molecule. Each Fab' fragment was separately secreted from E. coli and subjected to directed chemical coupling in vitro to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the ErbB2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets.

Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies have been produced using leucine zippers. Kostelny et al., <u>J. Immunol.</u> 148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers were reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology described by Hollinger et al., <u>Proc. Natl. Acad. Sci. USA</u> 90:6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain (V_H) connected to a light-chain variable domain (V_L) by a linker which is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L domains of one fragment are forced to pair with the complementary V_L and V_H domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See, Gruber et al., <u>J. Immunol.</u> 152:5368 (1994).

Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al., <u>J. Immunol.</u> 147:60 (1991). Exemplary bispecific antibodies can bind to two different epitopes, at least one of which originates in the protein antigen of the invention. Alternatively, an anti-antigenic arm of an immunoglobulin molecule can be combined with an arm which binds to a triggering molecule on

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a leukocyte such as a T-cell receptor molecule (e.g. CD2, CD3, CD28, or B7), or Fc receptors for IgG (FcγR), such as FcγRI (CD64), FcγRII (CD32) and FcγRIII (CD16) so as to focus cellular defense mechanisms to the cell expressing the particular antigen. Bispecific antibodies can also be used to direct cytotoxic agents to cells which express a particular antigen. These antibodies possess an antigen-binding arm and an arm which binds a cytotoxic agent or a radionuclide chelator, such as EOTUBE, DPTA, DOTA, or TETA. Another bispecific antibody of interest binds the protein antigen described herein and further binds tissue factor (TF).

5.13.6 Heteroconjugate Antibodies

Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (U.S. Patent No. 4,676,980), and for treatment of HIV infection (WO 91/00360; WO 92/200373; EP 03089). It is contemplated that the antibodies can be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins can be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptobutyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

5.13.7 Effector Function Engineering

It can be desirable to modify the antibody of the invention with respect to effector function, so as to enhance, *e.g.*, the effectiveness of the antibody in treating cancer. For example, cysteine residue(s) can be introduced into the Fc region, thereby allowing interchain disulfide bond formation in this region. The homodimeric antibody thus generated can have improved internalization capability and/or increased complement-mediated cell killing and antibody-dependent cellular cytotoxicity (ADCC). See Caron et al., J. Exp Med., 176: 1191-1195 (1992) and Shopes, J. Immunol., 148: 2918-2922 (1992). Homodimeric antibodies with enhanced antitumor activity can also be prepared using heterobifunctional cross-linkers as described in Wolff et al. Cancer Research, 53: 2560-2565 (1993). Alternatively, an antibody can be engineered that has dual Fc regions and can thereby have enhanced complement lysis and ADCC capabilities. See Stevenson et al., Anti-Cancer Drug Design, 3: 219-230 (1989).

5.13.8 Immunoconjugates

The invention also pertains to immunoconjugates comprising an antibody conjugated to a cytotoxic agent such as a chemotherapeutic agent, toxin (e.g., an enzymatically active toxin of

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bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive isotope (i.e., a radioconjugate).

Chemotherapeutic agents useful in the generation of such immunoconjugates have been described above. Enzymatically active toxins and fragments thereof that can be used include diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from Pseudomonas aeruginosa), ricin A chain, abrin A chain, modeccin A chain, alpha-sarcin, Aleurites fordii proteins, dianthin proteins, Phytolaca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, sapaonaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes. A variety of radionuclides are available for the production of radioconjugated antibodies. Examples include ²¹²Bi, ¹³¹I, ¹³¹In, ⁹⁰Y, and ¹⁸⁶Re.

Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutareldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as tolyene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta et al., Science, 238: 1098 (1987). Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See WO94/11026.

In another embodiment, the antibody can be conjugated to a "receptor" (such streptavidin) for utilization in tumor pretargeting wherein the antibody-receptor conjugate is administered to the patient, followed by removal of unbound conjugate from the circulation using a clearing agent and then administration of a "ligand" (e.g., avidin) that is in turn conjugated to a cytotoxic agent.

4.14 COMPUTER READABLE SEQUENCES

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media. A skilled

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artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially-available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g. text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

By providing any of the nucleotide sequences SEQ ID NO:1-1009 or a representative fragment thereof; or a nucleotide sequence at least 95% identical to any of the nucleotide sequences of SEQ ID NO:1-1009 in computer readable form, a skilled artisan can routinely access the sequence information for a variety of purposes. Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. The examples which follow demonstrate how software which implements the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990)) and BLAZE (Brutlag et al., Comp. Chem. 17:203-207 (1993)) search algorithms on a Sybase system is used to identify open reading frames (ORFs) within a nucleic acid sequence. Such ORFs may be protein encoding fragments and may be useful in producing commercially important proteins such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based systems are suitable for use in the present invention. As stated above, the computer-based systems of the present invention comprise a data storage means having stored

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therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means. As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of a known sequence which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, Smith-Waterman, MacPattern (EMBL), BLASTN and BLASTA (NPOLYPEPTIDEIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems. As used herein, a "target sequence" can be any nucleic acid or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 300 amino acids, more preferably from about 30 to 100 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzyme active sites and signal sequences. Nucleic acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

4.15 TRIPLE HELIX FORMATION

In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA.

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Polynucleotides suitable for use in these methods are preferably 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 15241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Olmno, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model

systems. Information contained in the sequences of the present invention is necessary for the

design of an antisense or triple helix oligonucleotide.

4.16 DIAGNOSTIC ASSAYS AND KITS

The present invention further provides methods to identify the presence or expression of one of the ORFs of the present invention, or homolog thereof, in a test sample, using a nucleic acid probe or antibodies of the present invention, optionally conjugated or otherwise associated with a suitable label.

In general, methods for detecting a polynucleotide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polynucleotide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polynucleotide of the invention is detected in the sample. Such methods can also comprise contacting a sample under stringent hybridization conditions with nucleic acid primers that anneal to a polynucleotide of the invention under such conditions, and amplifying annealed polynucleotides, so that if a polynucleotide is amplified, a polynucleotide of the invention is detected in the sample.

In general, methods for detecting a polypeptide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polypeptide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polypeptide of the invention is detected in the sample.

In detail, such methods comprise incubating a test sample with one or more of the antibodies or one or more of the nucleic acid probes of the present invention and assaying for binding of the nucleic acid probes or antibodies to components within the test sample.

Conditions for incubating a nucleic acid probe or antibody with a test sample vary. Incubation conditions depend on the format employed in the assay, the detection methods employed, and the type and nature of the nucleic acid probe or antibody used in the assay. One skilled in the art will recognize that any one of the commonly available hybridization.

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amplification or immunological assay formats can readily be adapted to employ the nucleic acid probes or antibodies of the present invention. Examples of such assays can be found in Chard, T., An Introduction to Radioimmunoassay and Related Techniques, Elsevier Science Publishers, Amsterdam, The Netherlands (1986); Bullock, G.R. et al., Techniques in Immunocytochemistry, Academic Press, Orlando, FL Vol. 1 (1982), Vol. 2 (1983), Vol. 3 (1985); Tijssen, P., Practice and Theory of immunoassays: Laboratory Techniques in Biochemistry and Molecular Biology, Elsevier Science Publishers, Amsterdam, The Netherlands (1985). The test samples of the present invention include cells, protein or membrane extracts of cells, or biological fluids such as sputum, blood, serum, plasma, or urine. The test sample used in the above-described method will vary based on the assay format, nature of the detection method and the tissues, cells or extracts used as the sample to be assayed. Methods for preparing protein extracts or membrane extracts of cells are well known in the art and can be readily be adapted in order to obtain a sample which is compatible with the system utilized.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the assays of the present invention. Specifically, the invention provides a compartment kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the probes or antibodies of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of a bound probe or antibody.

In detail, a compartment kit includes any kit in which reagents are contained in separate containers. Such containers include small glass containers, plastic containers or strips of plastic or paper. Such containers allows one to efficiently transfer reagents from one compartment to another compartment such that the samples and reagents are not cross-contaminated, and the agents or solutions of each container can be added in a quantitative fashion from one compartment to another. Such containers will include a container which will accept the test sample, a container which contains the antibodies used in the assay, containers which contain wash reagents (such as phosphate buffered saline, Tris-buffers, etc.), and containers which contain the reagents used to detect the bound antibody or probe. Types of detection reagents include labeled nucleic acid probes, labeled secondary antibodies, or in the alternative, if the primary antibody is labeled, the enzymatic, or antibody binding reagents which are capable of reacting with the labeled antibody. One skilled in the art will readily recognize that the disclosed probes and antibodies of the present invention can be readily incorporated into one of the established kit formats which are well known in the art.

4.17 MEDICAL IMAGING

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The novel polypeptides and binding partners of the invention are useful in medical imaging of sites expressing the molecules of the invention (*e.g.*, where the polypeptide of the invention is involved in the immune response, for imaging sites of inflammation or infection). See, *e.g.*, Kunkel et al., U.S. Pat. NO. 5,413,778. Such methods involve chemical attachment of a labeling or imaging agent, administration of the labeled polypeptide to a subject in a pharmaceutically acceptable carrier, and imaging the labeled polypeptide *in vivo* at the target site.

4.18 SCREENING ASSAYS

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Using the isolated proteins and polynucleotides of the invention, the present invention further provides methods of obtaining and identifying agents which bind to a polypeptide encoded by an ORF corresponding to any of the nucleotide sequences set forth in SEQ ID NO:1-1009, or bind to a specific domain of the polypeptide encoded by the nucleic acid. In detail, said method comprises the steps of:

- (a) contacting an agent with an isolated protein encoded by an ORF of the present invention, or nucleic acid of the invention; and
 - (b) determining whether the agent binds to said protein or said nucleic acid.

In general, therefore, such methods for identifying compounds that bind to a polynucleotide of the invention can comprise contacting a compound with a polynucleotide of the invention for a time sufficient to form a polynucleotide/compound complex, and detecting the complex, so that if a polynucleotide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Likewise, in general, therefore, such methods for identifying compounds that bind to a polypeptide of the invention can comprise contacting a compound with a polypeptide of the invention for a time sufficient to form a polypeptide/compound complex, and detecting the complex, so that if a polypeptide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Methods for identifying compounds that bind to a polypeptide of the invention can also comprise contacting a compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a receptor gene sequence in the cell, and detecting the complex by detecting reporter gene sequence expression, so that if a polypeptide/compound complex is detected, a compound that binds a polypeptide of the invention is identified.

Compounds identified via such methods can include compounds which modulate the activity of a polypeptide of the invention (that is, increase or decrease its activity, relative to

activity observed in the absence of the compound). Alternatively, compounds identified via such methods can include compounds which modulate the expression of a polynucleotide of the invention (that is, increase or decrease expression relative to expression levels observed in the absence of the compound). Compounds, such as compounds identified via the methods of the invention, can be tested using standard assays well known to those of skill in the art for their ability to modulate activity/expression.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention. Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like, capable of binding to a specific peptide sequence, in order to generate rationally designed antipeptide peptides, for example see Hurby et al., Application of Synthetic Peptides: Antisense Peptides," In Synthetic Peptides, A User's Guide, W.H. Freeman, NY (1992), pp. 289-307, and Kaspczak et al., Biochemistry 28:9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control. One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix formation by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods preferably contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription

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from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide and other DNA binding agents.

Agents which bind to a protein encoded by one of the ORFs of the present invention can be used as a diagnostic agent. Agents which bind to a protein encoded by one of the ORFs of the present invention can be formulated using known techniques to generate a pharmaceutical composition.

4.19 USE OF NUCLEIC ACIDS AS PROBES

Another aspect of the subject invention is to provide for polypeptide-specific nucleic acid hybridization probes capable of hybridizing with naturally occurring nucleotide sequences. The hybridization probes of the subject invention may be derived from any of the nucleotide sequences SEQ ID NO:1-1009. Because the corresponding gene is only expressed in a limited number of tissues, a hybridization probe derived from of any of the nucleotide sequences SEQ ID NO:1-1009 can be used as an indicator of the presence of RNA of cell type of such a tissue in a sample.

Any suitable hybridization technique can be employed, such as, for example, in situ hybridization. PCR as described in US Patents Nos. 4,683,195 and 4,965,188 provides additional uses for oligonucleotides based upon the nucleotide sequences. Such probes used in PCR may be of recombinant origin, may be chemically synthesized, or a mixture of both. The probe will comprise a discrete nucleotide sequence for the detection of identical sequences or a degenerate pool of possible sequences for identification of closely related genomic sequences.

Other means for producing specific hybridization probes for nucleic acids include the cloning of nucleic acid sequences into vectors for the production of mRNA probes. Such vectors are known in the art and are commercially available and may be used to synthesize RNA probes in vitro by means of the addition of the appropriate RNA polymerase as T7 or SP6 RNA polymerase and the appropriate radioactively labeled nucleotides. The nucleotide sequences may be used to construct hybridization probes for mapping their respective genomic sequences. The nucleotide sequence provided herein may be mapped to a chromosome or specific regions of a chromosome using well known genetic and/or chromosomal mapping techniques. These techniques include in situ hybridization, linkage analysis against known chromosomal markers, hybridization screening with libraries or flow-sorted chromosomal preparations specific to known chromosomes, and the like. The technique of fluorescent in situ hybridization of

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chromosome spreads has been described, among other places, in Verma et al (1988) Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York NY.

Fluorescent *in situ* hybridization of chromosomal preparations and other physical chromosome mapping techniques may be correlated with additional genetic map data. Examples of genetic map data can be found in the 1994 Genome Issue of Science (265:1981f). Correlation between the location of a nucleic acid on a physical chromosomal map and a specific disease (or predisposition to a specific disease) may help delimit the region of DNA associated with that genetic disease. The nucleotide sequences of the subject invention may be used to detect differences in gene sequences between normal, carrier or affected individuals.

4.20 PREPARATION OF SUPPORT BOUND OLIGONUCLEOTIDES

Oligonucleotides, *i.e.*, small nucleic acid segments, may be readily prepared by, for example, directly synthesizing the oligonucleotide by chemical means, as is commonly practiced using an automated oligonucleotide synthesizer.

Support bound oligonucleotides may be prepared by any of the methods known to those of skill in the art using any suitable support such as glass, polystyrene or Teflon. One strategy is to precisely spot oligonucleotides synthesized by standard synthesizers. Immobilization can be achieved using passive adsorption (Inouye & Hondo, (1990) J. Clin. Microbiol. 28(6) 1469-72); using UV light (Nagata *et al.*, 1985; Dahlen *et al.*, 1987; Morrissey & Collins, (1989) Mol. Cell Probes 3(2) 189-207) or by covalent binding of base modified DNA (Keller *et al.*, 1988; 1989); all references being specifically incorporated herein.

Another strategy that may be employed is the use of the strong biotin-streptavidin interaction as a linker. For example, Broude *et al.* (1994) Proc. Natl. Acad. Sci. USA 91(8) 3072-6, describe the use of biotinylated probes, although these are duplex probes, that are immobilized on streptavidin-coated magnetic beads. Streptavidin-coated beads may be purchased from Dynal, Oslo. Of course, this same linking chemistry is applicable to coating any surface with streptavidin. Biotinylated probes may be purchased from various sources, such as, *e.g.*, Operon Technologies (Alameda, CA).

Nunc Laboratories (Naperville, IL) is also selling suitable material that could be used. Nunc Laboratories have developed a method by which DNA can be covalently bound to the microwell surface termed Covalink NH. CovaLink NH is a polystyrene surface grafted with secondary amino groups (>NH) that serve as bridge-heads for further covalent coupling. CovaLink Modules may be purchased from Nunc Laboratories. DNA molecules may be bound to CovaLink exclusively at the 5'-end by a phosphoramidate bond, allowing immobilization of more than 1 pmol of DNA (Rasmussen et al., (1991) Anal. Biochem. 198(1) 138-42).

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The use of CovaLink NH strips for covalent binding of DNA molecules at the 5'-end has been described (Rasmussen et al., (1991). In this technology, a phosphoramidate bond is employed (Chu et al., (1983) Nucleic Acids Res. 11(8) 6513-29). This is beneficial as immobilization using only a single covalent bond is preferred. The phosphoramidate bond joins the DNA to the CovaLink NH secondary amino groups that are positioned at the end of spacer arms covalently grafted onto the polystyrene surface through a 2 nm long spacer arm. To link an oligonucleotide to CovaLink NH via an phosphoramidate bond, the oligonucleotide terminus must have a 5'-end phosphate group. It is, perhaps, even possible for biotin to be covalently bound to CovaLink and then streptavidin used to bind the probes.

More specifically, the linkage method includes dissolving DNA in water (7.5 ng/ul) and denaturing for 10 min. at 95°C and cooling on ice for 10 min. Ice-cold 0.1 M 1-methylimidazole, pH 7.0 (1-MeIm₇), is then added to a final concentration of 10 mM 1-MeIm₇. A ss DNA solution is then dispensed into CovaLink NH strips (75 ul/well) standing on ice.

Carbodiimide 0.2 M 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC), dissolved in 10 mM 1-MeIm₇, is made fresh and 25 ul added per well. The strips are incubated for 5 hours at 50°C. After incubation the strips are washed using, *e.g.*, Nunc-Immuno Wash; first the wells are washed 3 times, then they are soaked with washing solution for 5 min., and finally they are washed 3 times (where in the washing solution is 0.4 N NaOH, 0.25% SDS heated to 50°C).

It is contemplated that a further suitable method for use with the present invention is that described in PCT Patent Application WO 90/03382 (Southern & Maskos), incorporated herein by reference. This method of preparing an oligonucleotide bound to a support involves attaching a nucleoside 3'-reagent through the phosphate group by a covalent phosphodiester link to aliphatic hydroxyl groups carried by the support. The oligonucleotide is then synthesized on the supported nucleoside and protecting groups removed from the synthetic oligonucleotide chain under standard conditions that do not cleave the oligonucleotide from the support. Suitable reagents include nucleoside phosphoramidite and nucleoside hydrogen phosphorate.

An on-chip strategy for the preparation of DNA probe for the preparation of DNA probe arrays may be employed. For example, addressable laser-activated photodeprotection may be employed in the chemical synthesis of oligonucleotides directly on a glass surface, as described by Fodor *et al.* (1991) Science 251(4995) 767-73, incorporated herein by reference. Probes may also be immobilized on nylon supports as described by Van Ness *et al.* (1991) Nucleic Acids Res. 19(12) 3345-50; or linked to Teflon using the method of Duncan & Cavalier (1988) Anal. Biochem. 169(1) 104-8; all references being specifically incorporated herein.

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To link an oligonucleotide to a nylon support, as described by Van Ness *et al.* (1991), requires activation of the nylon surface via alkylation and selective activation of the 5'-amine of oligonucleotides with cyanuric chloride.

One particular way to prepare support bound oligonucleotides is to utilize the light-generated synthesis described by Pease *et al.*, (1994) PNAS USA 91(11) 5022-6, incorporated herein by reference). These authors used current photolithographic techniques to generate arrays of immobilized oligonucleotide probes (DNA chips). These methods, in which light is used to direct the synthesis of oligonucleotide probes in high-density, miniaturized arrays, utilize photolabile 5'-protected *N*-acyl-deoxynucleoside phosphoramidites, surface linker chemistry and versatile combinatorial synthesis strategies. A matrix of 256 spatially defined oligonucleotide probes may be generated in this manner.

4.21 PREPARATION OF NUCLEIC ACID FRAGMENTS

The nucleic acids may be obtained from any appropriate source, such as cDNAs, genomic DNA, chromosomal DNA, microdissected chromosome bands, cosmid or YAC inserts, and RNA, including mRNA without any amplification steps. For example, Sambrook *et al.* (1989) describes three protocols for the isolation of high molecular weight DNA from mammalian cells (p. 9.14-9.23).

DNA fragments may be prepared as clones in M13, plasmid or lambda vectors and/or prepared directly from genomic DNA or cDNA by PCR or other amplification methods. Samples may be prepared or dispensed in multiwell plates. About 100-1000 ng of DNA samples may be prepared in 2-500 ml of final volume.

The nucleic acids would then be fragmented by any of the methods known to those of skill in the art including, for example, using restriction enzymes as described at 9.24-9.28 of Sambrook *et al.* (1989), shearing by ultrasound and NaOH treatment.

Low pressure shearing is also appropriate, as described by Schriefer *et al.* (1990) Nucleic Acids Res. 18(24) 7455-6, incorporated herein by reference). In this method, DNA samples are passed through a small French pressure cell at a variety of low to intermediate pressures. A lever device allows controlled application of low to intermediate pressures to the cell. The results of these studies indicate that low-pressure shearing is a useful alternative to sonic and enzymatic DNA fragmentation methods.

One particularly suitable way for fragmenting DNA is contemplated to be that using the two base recognition endonuclease, *Cvi*JI, described by Fitzgerald *et al.* (1992) Nucleic Acids Res. 20(14) 3753-62. These authors described an approach for the rapid fragmentation and fractionation

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of DNA into particular sizes that they contemplated to be suitable for shotgun cloning and sequencing.

The restriction endonuclease *Cvi*JI normally cleaves the recognition sequence PuGCPy between the G and C to leave blunt ends. Atypical reaction conditions, which alter the specificity of this enzyme (*Cvi*JI**), yield a quasi-random distribution of DNA fragments form the small molecule pUC19 (2688 base pairs). Fitzgerald *et al.* (1992) quantitatively evaluated the randomness of this fragmentation strategy, using a *Cvi*JI** digest of pUC19 that was size fractionated by a rapid gel filtration method and directly ligated, without end repair, to a lac Z minus M13 cloning vector. Sequence analysis of 76 clones showed that *Cvi*JI** restricts pyGCPy and PuGCPu, in addition to PuGCPy sites, and that new sequence data is accumulated at a rate consistent with random fragmentation.

As reported in the literature, advantages of this approach compared to sonication and agarose gel fractionation include: smaller amounts of DNA are required (0.2-0.5 ug instead of 2-5 ug); and fewer steps are involved (no preligation, end repair, chemical extraction, or agarose gel electrophoresis and elution are needed

Irrespective of the manner in which the nucleic acid fragments are obtained or prepared, it is important to denature the DNA to give single stranded pieces available for hybridization. This is achieved by incubating the DNA solution for 2-5 minutes at 80-90°C. The solution is then cooled quickly to 2°C to prevent renaturation of the DNA fragments before they are contacted with the chip. Phosphate groups must also be removed from genomic DNA by methods known in the art.

4.22 PREPARATION OF DNA ARRAYS

Arrays may be prepared by spotting DNA samples on a support such as a nylon membrane. Spotting may be performed by using arrays of metal pins (the positions of which correspond to an array of wells in a microtiter plate) to repeated by transfer of about 20 nl of a DNA solution to a nylon membrane. By offset printing, a density of dots higher than the density of the wells is achieved. One to 25 dots may be accommodated in 1 mm², depending on the type of label used. By avoiding spotting in some preselected number of rows and columns, separate subsets (subarrays) may be formed. Samples in one subarray may be the same genomic segment of DNA (or the same gene) from different individuals, or may be different, overlapped genomic clones. Each of the subarrays may represent replica spotting of the same samples. In one example, a selected gene segment may be amplified from 64 patients. For each patient, the amplified gene segment may be in one 96-well plate (all 96 wells containing the same sample). A plate for each of the 64 patients is prepared. By using a 96-pin device, all samples may be spotted on one 8 x 12 cm membrane.

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Subarrays may contain 64 samples, one from each patient. Where the 96 subarrays are identical, the dot span may be 1 mm² and there may be a 1 mm space between subarrays.

Another approach is to use membranes or plates (available from NUNC, Naperville, Illinois) which may be partitioned by physical spacers *e.g.* a plastic grid molded over the membrane, the grid being similar to the sort of membrane applied to the bottom of multiwell plates, or hydrophobic strips. A fixed physical spacer is not preferred for imaging by exposure to flat phosphor-storage screens or x-ray films.

The present invention is illustrated in the following examples. Upon consideration of the present disclosure, one of skill in the art will appreciate that many other embodiments and variations may be made in the scope of the present invention. Accordingly, it is intended that the broader aspects of the present invention not be limited to the disclosure of the following examples. The present invention is not to be limited in scope by the exemplified embodiments which are intended as illustrations of single aspects of the invention, and compositions and methods which are functionally equivalent are within the scope of the invention. Indeed, numerous modifications and variations in the practice of the invention are expected to occur to those skilled in the art upon consideration of the present preferred embodiments. Consequently, the only limitations which should be placed upon the scope of the invention are those which appear in the appended claims.

All references cited within the body of the instant specification are hereby incorporated by reference in their entirety.

5.0 EXAMPLES

5.1 EXAMPLE 1

Novel Nucleic Acid Sequences Obtained From Various Libraries

A plurality of novel nucleic acids were obtained from cDNA libraries prepared from various human tissues and in some cases isolated from a genomic library derived from human chromosome using standard PCR, SBH sequence signature analysis and Sanger sequencing techniques. The inserts of the library were amplified with PCR using primers specific for the vector sequences which flank the inserts. Clones from cDNA libraries were spotted on nylon membrane filters and screened with oligonucleotide probes (e.g., 7-mers) to obtain signature sequences. The clones were clustered into groups of similar or identical sequences. Representative clones were selected for sequencing.

In some cases, the 5' sequence of the amplified inserts was then deduced using a typical Sanger sequencing protocol. PCR products were purified and subjected to fluorescent dye terminator cycle sequencing. Single pass gel sequencing was done using a 377 Applied Biosystems

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(ABI) sequencer to obtain the novel nucleic acid sequences. In some cases RACE (Random Amplification of cDNA Ends) was performed to further extend the sequence in the 5' direction.

5.2 EXAMPLE 2

Novel Contigs

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The novel contigs of the invention were assembled from sequences that were obtained from a cDNA library by methods described in Example 1 above, and in some cases sequences obtained from one or more public databases. Chromatograms were base called and assembled using a software suite from University of Washington, Seattle containing three applications designated PHRED, PHRAP, and CONSED. The sequences for the resulting nucleic acid contigs are designated as SEQ ID NO: 1-1009 and are provided in the attached Sequence Listing. The contigs were assembled using an EST sequence as a seed. Then a recursive algorithm was used to extend the seed EST into an extended assemblage, by pulling additional sequences from different databases (*i.e.*, Hyseq's database containing EST sequences, dbEST version 114, gb pri 114, and UniGene version 101) that belong to this assemblage. The algorithm terminated when there was no additional sequences from the above databases that would extend the assemblage. Inclusion of component sequences into the assemblage was based on a BLASTN hit to the extending assemblage with BLAST score greater than 300 and percent identity greater than 95%.

The nucleotide sequence within the assembled contigs that codes for signal peptide sequences and their cleavage sites was determined from using Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark). The process for identifying prokaryotic and eukaryotic signal peptides and their cleavage sites are also disclosed by Henrik Nielson, Jacob Engelbrecht, Soren Brunak, and Gunnar von Heijne in the publication "Identification of prokaryotic and eukaryotic signal peptides and prediction of their cleavage sites" Protein Engineering, vol. 10, no. 1, pp.1-6 (1997) incorporated herein by reference,. A maximum S score and a mean S score, as described in the Nielson et al. reference, are obtained from each assembled contig. Table 3 sets forth the nucleotide range for each sequence of SEQ ID NO: 1-1009 that encodes a corresponding amino acid sequence containing the signal peptide sequence and its cleavage site: the maximum S score and the mean S score obtained for each sequence.

A signal peptide or leader peptide is usually a segment of about 15 to 30 amino acids at the N terminus of protein that enables the protein to be targeted to a cell membrane or secreted from a cell. Generally, the signal peptide acts as an export lable and is removed as the protein is secreted in its final form.

The nearest neighbor result for the assembled contig was obtained by a BLASTX version 2.01al 19 MP-Washington University search against Genpept release 120 and Geneseq database (October 12, 2000, update 21 (Derwent)), using BLAST algorithm. The nearest neighbor result showed the closest homologue for each assemblage from Genpept (and contains the translated amino acid sequences for which the assemblage encodes). The nearest neighbor results for SEQ ID NO: 1-1009 are shown in Table 2.

Tables 1, 2 and 3 follow. Table 1 shows the various tissue sources of SEQ ID NO: 1-1009. Table 2 shows the nearest neighbor result for the assembled contig. The nearest neighbor result shows the closest homolog with an identifiable function for each assemblage. Table 3 contains the start and stop nucleotides for the translated amino acid sequence for which each assemblage encodes. Table 3 also provides a correlation between the amino acid sequences set forth in the Sequence Listing, the nucleotide sequences set forth in the Sequence Listing and the SEQ ID NO. in USSN 09/491,404.

15

10

TABLE 1

169 171-172 175-176 181 203 212 202 223 221-252 258 263 267 279 36 343 388 396 400-401 422 212 428-429 431 437 456 464 487 503 513 524 561 500 588 609 619 682 812 946 598 965 980 983 989 999 812 946 598 965 980 983 989 999 87 111-113 116 122-123 129 139 143 148 159 163 167 175-176 178 181 183 186 210-204 206 208-209 212 214 220 222 228 230 234-235 237 246 249-250 252 255 255 259 262- 264 266-267 279-280 286 329 336 351 388 379 396 422 429 431 437 439 444-445 450 452 456 467-468 479 444 504 50 452 456 467-468 479 444 504 50 452 456 467-468 479 444 504 50 452 456 467-468 479 444 50 450 452 456 467-468 486 588 590 902 906 910 922-924 332-933 941 945 955 956 1-562 578 86 580 583 550 563 559 561-562 578 876 885 590 902 906 910 922-924 332-933 941 945 955 958 965 971 983-984 999 990 100 adult brain Clontech ABR001 81 122 148 181 183 204 207 233 237 250 267 301 346 394 396 437 439 457 505 563 618 653 655 721 764 769 775 885 92 949 adult brain Clontech ABR006 148 152 222 277 269 583 640 677 878 adult brain Clontech ABR008 2 10-11 3-14 19-20 23 28-29 344 35 37 39-40 45 49-50 52 60 73-74 78 88 791 94 98 101 109 114-117 122-123 143 145 146 149-15 152 156 162 168 173-178 181 183 187 189 194 20 206-209 212 214-215 220- 221 228 231 233-238 246-247 249- 253 255-260 262 266 269-70 272 276 278-281 284 294 301 313 3167 189 253 255-260 262 266 269-70 272 276 278-281 284 294 301 313 3167 189 38 390-392 396 400-401 403 405- 407 414 417 427-423 425 427-428 433 437 441 443-446 452-453 456 464 467 469 473-479 489 448 487- 488 491 497-498 500 502 504-505 507 719-505 595 595 595 595 595 595 595 595 595	TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY NAME	SEQ ID NOS: OF NUCLEOTIDE(S)
87 111-13 116 122-123 129 139 13 148 189 163 167 172-716 178 131 183 166 201-204 206 208-209 212 214 220 222 228 230 234-225 227 246 249-250 252 255 259 262- 264 266-267 279-280 286 329 336 351 358 379 396 422 429 431 437 439 444-445 450 452 455 467-468 479 484 503-504 507 135 523-524 526 533 550 553 559 561-562 578 580 583 636 636 640 680 711 759 764 769 772 799 803 824 830 842 865 885 900 902 906 901 922-924 932-933 941 945 951 955 958 965 971 983-984 989 999 1005 adult brain Clontech ABR001 81 122 148 181 183 204 207 233 277 250 267 301 346 394 396 437 439 457 505 563 618 653 655 721 764 795 885 942 249 adult brain Clontech ABR008 2 10-11 13-14 19-20 23 28-29 34- 35 37 39-40 45 49-50 560 73-74 78 83 87-91 94 98 101 109 114-117 122-123 143 145 148-150 152 156 162 168 173-178 181 181 187 189 194 204 206-209 212 214-215 220- 221 228 231 233-238 246-247 249- 233 255-260 262 266 269-270 272 2765 278-281 284 294 301 313 316- 330 335 337-338 343 363 372 379 388 390-392 396 400-401 403 405- 407 414 417 422-423 425 427-428 433 437 441 442-446 45-437 456- 488 491 497-498 500 504-505 507 519-520 521-526 533 544-545 553 555-556 563 570-571 574-576 578-580 583 615 618-619 637-638 643-644 653 655-656 661 663 678 680 689-690 695 699 702 705 717- 718 720-727-77-77 784-796 500 504-508 680 689-690 695 699 702 705 717- 718 720-727-77-77 78 78-786 588 888 890-999 500-900 910 922 999 500-900 900-900 900-900 900-900 900-900-900 900-900 910 912 920 930-932 939- 942 946-947 951-955 955 956 966 987-989 999 9900 995 999 9900-900 940 980 9819 8812 814-815 817 834 840 842 844-846 852 855-856 888-860 870-873 875 877 885-886 888-880- 887 989 999 999 999 999 995 999 999 999 999	adult brain	GIBCO .	AB3001	169 171-172 175-176 181 203 212 220 222 230 251-252 258 263 267 279 336 343 358 396 400-401 422 428-429 431 437 456 464 487 503 513 524 561 580 583 609 619 682
237 250 267 301 346 394 396 437 439 457 505 563 618 653 655 721 764 795 885 942 949 adult brain	adult brain		ABD003	5 23 26 28-29 31 34-36 61 74 78 87 111-113 116 122-123 129 139 143 148 159 163 167 175-176 178 181 183 186 201-204 206 208-209 212 214 220 222 228 230 234-235 237 246 249-250 252 255 259 262- 264 266-267 279-280 286 329 336 351 358 379 396 422 429 431 437 439 444-445 450 452 456 467-468 479 484 503-504 507 513 523-524 526 533 550 553 559 561-562 578 580 583 636 638 640 683 711 759 764 769 772 799 803 824 830 842 865 885 900 902 906 910 922-924 932-933 941 945 951 955 958 965 971 983-984 989 999 1005
## ABROOS 2 10-11 13-14 19-20 23 28-29 34-35 37 39-40 45 49-50 52 60 73-74 78 83 87-91 94 98 101 109 114-117 122-123 143 145 148-150 152 156 162 168 173-178 181 183 187 189 194 204 206-209 212 214-215 220-221 228 231 233-238 246-247 249-253 255-260 262 266 269-270 272 276 278-281 284 294 301 313 316-320 335 337-338 343 363 372 379 388 390-392 396 400-401 403 405-407 414 417 422-423 425 427-428 433 437 441 447-448-452-453 456 464 467 469 473-479 482 484 487-488 491 497-498 500 502 504-505 507 519-520 523-526 563 570-571 574-576 578-580 583 615 618-619 637-638 643-644 653 655-656 661 663 678 680 689-690 695 699 702 705 717-718 720 722 725-726 742 746 752 754-755 759 761 763-765 767 769 772-774 776 784-789 792 795 799 809-810 812 814-815 817 834 840 842 844-846 852 855-856 888 890-897 903-904 910 928 930-932 939-942 946-947 951-952 955 957 960 964-965 679 711 975-976 978 986-997 997 997 999 999 1001	adult brain	Clontech	ABR001	237 250 267 301 346 394 396 437 439 457 505 563 618 653 655 721
35 37 39-40 45 49-50 52 60 73-74 78 83 87-91 94 98 101 109 114-117 122-123 143 145 148-150 152 156 162 168 173-178 181 183 187 189 194 204 206-209 212 214-215 220- 221 228 231 233-238 246-247 249- 253 255-260 262 266 269-270 272 276 278-281 284 294 301 313 316- 320 335 337-338 343 363 372 379 388 390-392 396 400-401 403 405- 407 414 417 422-423 425 427-428 433 437 441 443-446 452-453 456 464 467 469 473-479 482 484 487- 488 491 497-498 500 502 504-505 507 519-520 523-526 533 544-545 553 555-556 563 570-571 574-576 578-580 583 615 618-619 637-638 643-644 653 655-656 661 663 678 680 689-690 695 699 702 705 717- 718 720 722 725-726 742 746 752 754-755 759 761 763-765 767 769 772-774 776 784-789 792 795 799 809-810 812 814-815 817 834 840 842 844-846 852 855-856 888 890- 897 903-904 910 928 930-932 939- 942 946-947 951-952 955 957 960 964-965 967 971 975-976 978 986- 987 989 992 999 1001	adult brain	Clontech	ABR006	
987 989 992 999 1001	dure brain	CIONICECII	ADRUUG	35 37 39-40 45 49-50 52 60 73-74 78 83 87-91 94 98 101 109 114-117 122-123 143 145 148-150 152 156 162 168 173-178 181 183 187 189 194 204 206-209 212 214-215 220- 221 228 231 233-238 246-247 249- 253 255-260 262 266 269-270 272 276 278-281 284 294 301 313 316- 320 335 337-338 343 363 372 379 388 390-392 396 400-401 403 405- 407 414 417 422-423 425 427-428 433 437 441 443-446 452-453 456 464 467 469 473-479 482 484 487- 488 491 497-498 500 502 504-505 507 519-520 523-526 533 544-545 553 555-556 563 570-571 574-576 578-580 583 615 618-619 637-638 643-644 653 655-656 661 663 678 680 689-690 695 699 702 705 717- 718 720 722 725-726 742 746 752 754-755 759 761 763-765 767 769 772-774 776 784-789 792 795 799 809-810 812 814-815 817 834 840 842 844-846 852 855-856 858-860 870-873 875 877 885-886 888 890- 897 903-904 910 928 930-932 939- 942 946-947 951-952 955 957 960
	adult brain	Clontech	ABR011	987 989 992 999 1001 214 965

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY NAME	SEQ ID NOS: OF NUCLEOTIDE(S)
adult brain	BioChain	ABR012	152 498
adult brain	Invitrogen	ABR013	142 207 254 396 442 498
adult brain	Invitrogen	ABT004	2 23 31 34 78 96 116 129 141 160 176-177 181 183 202 214 231 233 248 256 258-260 262 278 310 336- 337 379 416 437 439 443-444 450 452 454 464 467 479 484 500 504
			519 526 553 570 590 619 638 640 647 653 655 678 711 759 764 789 795 799 885 887 892 902 905 907 910 915 922 941-942 955 960 989 999
cultured	Strategene	ADP001	17 37 39 74 79 111 129 152 160
preadipocytes			200 222 248 252 268 274 358 385 450 456 504 526 571 583 619 633 640 740 803 816 829 842 887 939- 940 965 973 977 986
adrenal gland	Clontech	ADR002	4 6 19 36 39 49 51-53 74 76 118 122-123 147-148 152 156 160 167 171-172 181 183 204 206 212 223- 224 228 233-234 246 249-250 254- 255 262 274 278-279 284 287 294 317 336 355 358 366 379 392 401-
			402 412 417 420 431-432 439 464 470 479-480 484 503-504 506 509 519 524 526-527 541 553 555 561 583 614 619 631 638 646 682 738- 739 756 760 764 770 800 802-803 816-817 838 847 852 863 881 887 905-906 910 923 926 932 941 950-
			951 989 999 1002
adult heart	GIBCO	AHROOI	6 20 26 29 31 34 37 39 41 46 61 74 78 101 114 116-118 122-124 128 145 147-148 152 155 163 175-176 178 181 183 200 204 206 210 212 215 228 230 234-235 237 246 248- 252 255-256 262-263 266-268 272 278 280 282-283 286 294 309 313 350-351 358 370 374 379 391-392 394 397 400-401 409 420 423 431- 432 434 436 438 441 443 452 455- 456 461 467-468 479-480 484 487 498 500 503 505 511 519 533 541 550 552-553 558 561-562 568 575 583 590 597-598 603 619 636-638 644-645 667-668 680 684 711-712 714-715 723 732 750 789 803 805 816 822 828 885 889 900 902 905 908 910 916-917 923-924 932 935 937 939 941 950 952 954 960 965 974 982 984 987 993 1005
adult kidnov	GTRCO	AKD001	4 13-14 19-20 23 26-31 37 39 47
adult kidney	GIBCO	ARDUUI	4 13-14 19-20 23 26-31 37 39 47 49 54 61 64 78 81 87 91 98 101 114 118 122-123 127 129-130 141- 143 145 148-149 155-158 160 163 168 171-172 175-176 178-181 183 197-198 200 203-206 208 212 215 221-222 228 230 234 237 241 245- 246 250-252 254-257 262-263 265- 269 278-279 282-284 286 297 301

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY	SEQ ID NOS: OF NUCLEOTIDE(S)
		NAME	308 333 336 352-353 358 371-372 379 381 386 391 394 396-397 400-
			401 405 409 417 420 428-429 431
			436-437 443 445 450 456 463-466
			468 475 479-480 484 487 495 498-
			499 503-505 507 511 513 517 523
			526 529 533 539 541-542 550 552-
			553 555 561 570-572 575 577-578
			583 587 597 604 606 609 619 636
			638 640-642 648 680 682 701 706
			714 721 732 740 747 771 792 803 805 809 811-812 829 838 842 862
			865 885 889 900 902 905-906 908
			910-911 918-921 924 926 928-930
			937 939 941-942 950-951 953 955
			958 960 963 965 967 976 978-979
			982-984 1005
adult kidney	Invitrogen	AKT002	19 31 78 81 91 98-99 122 142 145
_			148 152 158 169 176 248 254 256
			262 266 279 296-297 301 321 353
			372 401 405 416 420 429-430 441
			456 464 498 504 507 523 526 533
			541 583 592-597 649 701 791 838
			862 868 911 926 933 946-947 958
2917 + 71122	GIBCO	ALG001	960 971
adult lung	GIBCO	ALGUUI	145 148 179 183 194 198 200 205
			212 220 228 234 246 248 250-251
			254-255 263 268 277 279 289 298
			306 337 343 372 379-380 385 401
			405-406 408 410 420 431 440 443
			445 449 455 484 499 503 507 513
	•		517 571 590 597 617 636 640 714
			732 749-750 805 885 900 905 910
			918 941 955 958 960 977 980 1001
			1005
lymph node	Clontech	ALN001	43 48 53 108 123 136 142 147 160
		•	178 181 183 200 205 228 244 246 250 254 268 270 291 379 399 419
			431 440 442 479-480 484 519 533
			539 553 559 565 583 616-617 619
			636 662 701 740 805 833 910 913
			928 941 977
young liver	GIBCO	ALV001	19 42 45 61 64 84 98 107 109 122-
			123 129-130 133 142 148 168-169
			178 181 183 200 205 207 227-229
			232 238 246-248 250 253-255 262-
			263 265 268 279 317 336 371 377
			392 400 410 431 436-437 443 445
			448-450 484 487 513 533 545 559
			561 570 578 617 632 638 640 648 680 771 803 816 836-838 885 906
			926 940 986
adult liver	Invitrogen	ALV002	13-14 26 36 54 64 74 76 109 117
addic Tivel	1111110109CH	ABVOUZ	122 179 181 183 187 204 215 221
			225 229 232 247-248 250 256-257
			275 304 307 315 317 321-322 371
			377 379 386 416 420 448-449 457
			464 475 479 481 483-484 504 507
		1	526 553 557 570 619 627-629 632

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEO	SEQ ID NOS: OF NUCLEOTIDE(S)
TISSUE ORIGIN	RNA SOURCE	LIBRARY	SEQ ID NOS. OF NOCHEOTIDE(S)
		NAME	
		NAI-III	638 640 653 655 675 680 701 752
			768 827 848 865 882 885 889 910
			951 955 959 963 967 978 989 999-
			1000
adult ovary	Invitrogen	AOV001	4 12 19 23 28-32 34-37 39 45 48
			52 54 60-61 64-65 67 76 78 87 96
			98-100 108 111-112 114 116-118
			122-123 126 129-130 132-134 137
			139 142-145 147-149 152 162-163
			169-172 176 178 180-183 187 191-
			192 197-202 204-206 212 214-217
			219-222 228 234-235 237 242 246-
			248 250-252 254-256 262 265-269
			274 279-280 282-284 294 308-309
			313 317 336-337 346 358 361 364
			371 374 379 391-392 394 396-397
		1	400 408 414 418 420 423 425 428-
			429 431 435-437 440-441 443-447
			450 452 455-459 463-464 467-468
			479-480 484 487 492 495 499-500
			503 505 512-513 517 519 524 533
			539 545 553 555 557-559 561 565-
			566 568 571 575 577-578 581 583
			590 597 605 610 613 616-617 619
			636 638 640 645-646 649-650 654
			662 671 680 682 694 697 701 711
1			732 735 739-741 750 753 760 764
			771 780 785 789 792 803 806 810
			812 821 831-832 838 841-842 879
			885 887 900 902 905-906 908-912
			917 921-922 924 928 936-939 941-
			942 946 950-952 957-958 960 962-
			965 979 982 987 989 994 998-999
			1005 1008
adult placenta	Clontech	APL001	122 148 168 181 194 200 248 262
			268 317 436 541 561 803 838 911
			971
placenta	Invitrogen	APL002	38 61 78-79 142 149 176 187 194
1			206 215 246 252 278 337 346 379
			400 456 464 478-479 484 487 504
			519 526 553 571 638 640 732 842
			910-911 918 941 958
adult spleen	GIBCO	ASP001	23 26 39 43 48 61 63 78 87 98 108
addit Spicen	CIDCO	7101001	110 123 136 142 157 176 178 181
			183 197-198 201-202 205-206 213
			220 222 228 234 237 244 250-252
			254-255 257 263 294 305 320 336-
			337 354 358 371-372 376 379 397
			400 405 410 414 431 437 440 455-
			456 484 487 498-499 504 506-507
			511-512 519 523 526 529 533 539
			550 561 565 572 575 583 586 597
			616-617 619 621 636 640 687 701
			713 732 740 748 803 812 816 835
			910 930 939 946 956 958
testis	GIBCO	ATS001	20 23 29 61 64 76 114 123 126 143
			145 148-149 175 178 182 200 203
			206 209 235 248 252 257 263 268
			279-281 283-284 333 358 371 391
			396 400 418 423 431 438-439 441
L			1 220 400 440 450 431 430-432 447

TABLE 1

TISSUE ORIGIN	RNA SOURCE	Trycho	SEO ID NOS: OF NUCLEOTIDE(S)
TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY	SEQ ID NOS: OF NOCLEOTIDE(S)
		NAME	
		MAME	445 456 479-480 487 490 505 507-
			508 516-517 521 524 533 550 559
			561-562 582 597 606 638 646 676
			680 750 772 803 834 877 908 911
			914 937-938 950 989 999
adult bladder	Invitrogen	BLD001	23 37 77-78 84 160 176 178 181
addic bradder	invictogen	BIDOOL	215 218 248 252 262 274 299 334
]		351 401 464 474 484 517 543 619
			663 692 729 908 910 918 937 941
			951 960 962
bone marrow	Clontech	BMD001	19 31 39 43 48 52-53 95-96 98 100
Done marrow	Cronecen	DIADOUT	108 111-112 114 117 122-123 136
			141-142 144-145 147-149 152 161
			163 169 181 183 187 194 201 204-
			205 208 213 222 228 234 241-242
			244-246 248-251 254-255 257 267
	1		272 274 282 286 288-289 292 294
			313 317 335 337 339 346-347 358
	-		
			363 365 374 379 391-392 395-398
			406 408 414 418 423 428 436 440-
	1		498-500 504 508 511 516 519 526 533 539 541 553 556 559 561 565
			571 573 583 597 612 617 619 638
	i		640 646 649 651 677 681 685 707
			709-710 721 734 764 771 803 806
			811 838 852 858 869 885 908 910
			916 922 930 936-937 941 951 965
1	G1 and a sh	DMD000	982 985 989 991 995 999 1005 1008
bone marrow	Clontech	BMD002	31 39 43 48 68 71 91 108 122-123 134 136 142 148-150 152 161 169
	1		· · · -
			178 181 194 196 204-205 208 244
	İ		246 254 262-263 265 267 272-273
			300 320 343 356 363 372 379 405
			408 413-414 430-431 436 440-441 454 479 484 486 512-513 517 519
			533 553 559 570 583 590 617-619
			634 637 651 674 692 793-794 800
1			803 818 852 880 904 910 930 936
			941 950
hono marrow	Clentech	TMD004	142 152 254 274
bone marrow adult colon	Clontech	BMD004	
adult colon	Invitrogen	CLN001	26 29 48 61 108-109 129-130 144
			176 194 215 221 252 401 436 440
			450 498 511 533 583 590 616-617
adult corrie	PioChain	CVVCC1	706 764 905 939 955
adult cervix	BioChain	CVX001	6 16 19-20 29 35 37 43 45 64 73
			75-76 86 92 96-98 100-101 105 108
			111 113 122 143 145 147-149 163-
			165 167 172 174 178 181-183 187
			200-201 206 222 234 237-238 242-
			243 246 248 250-251 253 261-262
			265 268 270 274 279 283-284 294
			308 343 345 352 365 379 381 391
			400 409 420 423-424 428 436 443-
			444 463-464 473 479-480 484 487
			505 508 510-512 516-517 519 523-
		1	1 ED4 EDD EDD EED EED EED EC1
		į	524 533 539 553-555 558-559 561-
			562 575 578 583 591 597 619 643
			•

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ	SEQ ID NOS: OF NUCLEOTIDE(S)
		LIBRARY NAME	
			910 926-927 933 937 941 960 963
			965 967-968 977 982 989 999 1008-
			1009
diaphragm endothelial	BioChain	DIA002	26 152 499 680
cells	Strategene	EDT001	13-14 19 23 26 30-32 34 39 67 73- 74 76 78 91 101 109 114 116 118
00115			129 145 149 152 156 160-161 167
			176 180 183 187 197 201 203-204
			206 209 215 222 226 228 230 237
			246 248 250-252 256-257 262 266
			276 279 282-283 286 309 312-313
			343 358 372 391-392 394 396 400-
•			401 405 409 413 420 423 429-431 436 438 443-445 450 455-456 479
			484 487 498-499 503 507 509 511
			513 523 561-562 571 575 583 619
			639 646 653 655 680 711 721 729
			739 771-772 775 779 795 803 805
			834 838-840 885 889 900 905-906
			911 917-918 922 924 930 942 946
Genomic clones	Genomic DNA	EPM001	955 958 960 977-979 982-984 122 148 436
from the short	from Genetic	EFMOOT	122 140 430
arm of	Research		
chromosome 8			
Genomic clones	Genomic DNA	EPM003	122 148 379 436
from the short	from Genetic		
arm of	Research		
chromosome 8 Genomic clones	Genomic DNA	EPM004	122 148 436
from the short	from Genetic	BPNO04	122 140 430
arm of	Research		
chromosome 8			
Genomic clones	Genomic DNA	EPM005	148
from the short	from Genetic		
arm of chromosome 8	Research		
esophagus	BioChain	ESO002	152 178 583
fetal brain	Clontech	FBR001	122 148 181 279 284 484 553 575
	0101100011	1 DROUL	619 668 911
fetal brain	Clontech	FBR004	122 190 212 379 479 484 541 905
			922 924 941 950
fetal brain	Clontech	FBR006	2 23 31 36 39 42 44 49 52 78 87
			114 117 122-123 145 148 176-177
			180-181 187 204 208 210 215 220 235 238-239 241 245-246 251 253
			256 259 266 270 278 280 286 314
			317 337 372 379 392 396 400-401
			405-406 410 414 423 428 439-440
			443 445 452 467 473 479 484 487
			491 497 500 504 517 519 524 526
			544 553 556 561 563 568 570-571
			573 577 586 619 647 653 655 664-
	İ		665 680 739 742 746 754 766 772- 776 784 795 798 834 840 842 863
			878 885 892-893 898-899 910 930
			941-942 946 952 965 971 976 987
			993
fetal brain	Invitrogen	FBT002	19 31 34-35 44-45 78-79 87 96 101
			116 129 176 181 204 206 233 235

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ	SEQ ID NOS: OF NUCLEOTIDE(S)
		LIBRARY	
		NAME	
			256-257 259 262 278 280 317 320
			337 380 396-397 401 437 443 446
			450 453 464 480 484 498-499 504
			526 577 591 619 640 664 680 697
fetal heart	Tarritanoan	FHR001	710 764 900 902 905 910 958 500 910
fetal kidney	Invitrogen Clontech	FKD001	39 47 96 98 122-123 148 156 181
recar kruney	CIOntech	FKDUUI	200 207 246 268 274 279 283 300
			379 411 445 464 468 479 484 506
			542 553 561 583 619 680 686 712
			747 910 941
fetal kidney	Clontech	FKD002	479 484 583 803 910 941
fetal kidney	Invitrogen	FKD007	864
fetal lung	Clontech	FLG001	64 96 143-144 168 194 206 234 266
_			335 337 363 500 507 561 619 968
fetal lung	Invitrogen	FLG003	3 13-14 55 61 79 122-123 148 160
			181 183 194 200 234 248 250 252
			266 268 273 289 294 336 358 428
			432 436 484 507 510 513-514 533
			541 557-558 582-583 597 671 711
	71	TT 0004	764 777 806 811 817 905 933 978
fetal lung fetal liver-	Clontech Columbia	FLG004 FLS001	951
spleen	University	FLSOUI	13-15 19-21 23-26 28-30 32 34 37 39 45 47-49 56 67 72-74 78 84 87
spreen	Oniversity		91 96-98 101 103-104 108 111 114
			116 122-123 126 129 131 133 142-
			145 147-149 151-152 156 160-161
			166 168-169 172 176 178-179 181
			183-185 192-194 197-202 204-206
			208 215 221-222 224 228-229 232
			234-235 237 246 248-252 254-257
			262 266-268 272 274 278-280 282-
			287 294 313 315 321 333 336-337
			343-344 358 372 377-379 386 391-
			393 397 400-402 404-405 409-410
			418 420-421 429 431 436-437 440-
			441 443 445 448-450 456-457 464
			473 475 478-481 483-484 487-488
			498 500 503 505 507 509 513 522-
			523 528 533-534 541 551 553 558
			560-562 564-565 570 575 577-578
			583 586 590 597 600 605-607 617
			619 632 636 638 640 644 646 672
			677-680 705 711 729 732 735-738 740 742 748 760 763-764 771-772
			792 802-803 805-806 812 816-817
			820-821 824-827 834 838 842-843
			848 853 861 865 878 885 887 889
			900 902 904-906 908 910-911 917
			924 926 928 930 934 936-937 941
		1	944 946 950-951 955 958 960 963
			965 974-980 982-983 988-990 999
fetal liver-	Columbia	FLS002	4 8 12 15-16 18-21 23-24 26 32 37
spleen	University		39 47 54 61 64 67 71-72 74 76 79
-			83-84 87 91 96-98 100-104 109
			111-113 122-123 129 133 141 145
			147-149 152 161 163 169 171-172
			174 178-181 183 185 187-188 192-
	,	1	
		ĺ	195 198-202 205 207-209 213 215 221-222 229 232 234-235 237 241

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ	SEQ ID NOS: OF NUCLEOTIDE(S)
TIBBOE ORIGIN	KNA SOURCE	LIBRARY	SEQ ID NOS. OF NOCEBOTIDE (3)
		NAME	1
			244-246 248 250 262 265 267-268
			270 274 278-280 283-284 290 294
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			346 351-352 358 360-361 371-372
			377 382 391-393 397 399-401 404-
			405 410 414 425 429 431 436 440-
			441 445-446 448-450 453 456 464
			473 475 479-480 487 492 498 500
			503-504 507 512 517 519 523 526
			540 557 561-563 565 574-575 577-
			578 583 590 597 605-606 608 611
			614 616 619 631-634 636-638 640
			646 649-650 662 671-673 676-678
			682 684 701-702 704-705 711 716
			732 735 748 760 762-764 768 771-
†			772 779 790 802 805 815-816 834
			838 842 848 865 878-879 883 887-
			889 903 905-906 910 916-917 922
			924 928 930 939 944 946 950 955-
1			956 958 960 965 975 977 982-983
		77.0000	987-988 993-994 998 1004
fetal liver-	Columbia	FLS003	377 732 889 938
spleen	University	FLV001	22 20 20 04 100 104 200 221 222
fetal liver	Invitrogen	LTAGGI	23 29 39 84 109 194 208 221 232 247-248 278 301 321 336-337 370-
			371 379 443 448-449 464 475 479-
			480 498 500 533 550 578 590 632
			636 640 678 680 683 751 763 803
			882-883 885 887-889 910 921 942
			946 951 963 988
fetal liver	Clontech	FLV004	37 122 200 232 268 274 377 583
Tecar Trver	Cronceen	15004	946
fetal muscle	Invitrogen	FMS001	29 37 41 64 66 74 148 164 200 202
			208-209 252 257 259 262 265 268
			274 279 337 346 379 445 480-481
			505 507 553 555 561 571 606 640
			676 781 801 838 910 926 928 951
			957 960 963 965
fetal muscle	Invitrogen	FMS002	200 268 274
fetal skin	Invitrogen	FSK001	23 29 31 34 49 78 84 87 96 100
	_		112 116 133 143 148 163 168 172
			176-177 181 193 199-202 208 215
1			222 235 240 246 248 252 256-257
			262-268 274 280 282 294 309 314
			317 322 346 358 371 373-375 379
			414 417 419-420 436-437 441 445
			454 456 458 479-480 484 499-500
			504 507 513 519-520 526 533 539
			541 545-547 550 561 565 570-571
			575 577 583 590 598-599 619 644
			650 665 697 702 706 739 742 744
]	1		784 790 792-793 812 816 861 877
			889 906 910 918 922 941 949 951-
			952 955 962 964-965 968 979 983
			987 989 999
fetal skin	Invitrogen	FSK002	200 257 265 268 274 513 688
	T 70 (10)	FSP001	1 30 431 503 533 617
fetal spleen	BioChain		39 431 523 533 617
fetal spleen umbilical cord	BioChain	FUC001	19 28-29 34 39 74 96 99 101 111

TABLE 1

	Ditt. COTTLOT	111000	CDO TO MOS OF MISSIFICATION
TISSUE ORIGIN	RNA SOURCE	HYSEQ	SEQ ID NOS: OF NUCLEOTIDE(S)
		LIBRARY	
		NAME	
			222 228 230 237-238 246 248 252-
			253 255 257 259 262 265 268-269
	İ		272 274 282 325 351 379 396 400-
			401 413 429 441 443 445 452 456-
	•		457 467-468 479 484 487 505 513
			517 519 523 533 541 553 555 561
	1		571 575 577 583 590 601-602 605-
	i		606 619 636 645 680 693 698 711
			757 759 764 803 814 816 821 853
			885 889 900 906 908 910 924 926
			932 937 941 943 946 951-952 955
			958 976 987 989 993-994 999
£24-3 13	GERGO	TIPDOOT	
fetal brain	GIBCO	HFB001	13-14 19 26 29 31-32 39 44-45 61
			67 74 78 88 100 114 122-123 126
			129 148 152 163 167 169 171-172
			175-176 180-181 187 201-204 206
			209 212 215 220 222 227-228 230
			233-235 237 246 249 251 258-259
			262-263 266 269 279-280 282 284
			286 333 337 340 342 355 358 362
			366 379 391 394-397 406 422-423
			428-429 431 436-437 443-446 450
			452 456 467-468 479-480 484 498
			504-505 513 517 523 526-527 533
		\	539 541 558-559 561-562 574 580
			583 605 619 635 638 643 680 682
			708 711 739-740 742 764 776 803
			812 823 865 885 900 902 905 910
			0.00 0.00 0.00 0.00 0.00 0.00
			917 924 928 932 939 941 945 958
			960 964-965 974 978-979 984
macrophage	Invitrogen	HMP001	960 964-965 974 978-979 984 152 201 498 983
macrophage infant brain	Invitrogen Columbia	НМР001 IB2002	960 964-965 974 978-979 984
			960 964-965 974 978-979 984 152 201 498 983
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255-
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467-
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958
	Columbia		960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500 504 526 575 583 590 605 630 653
infant brain	Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500 504 526 575 583 590 605 630 653 655 703 733 757 764 795 865 884-
infant brain	Columbia University Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500 504 526 575 583 590 605 630 653 655 703 733 757 764 795 865 884- 885 900 905 919 924 974-975 982
infant brain	Columbia University Columbia University Columbia	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500 504 526 575 583 590 605 630 653 655 703 733 757 764 795 865 884-
infant brain	Columbia University Columbia University	IB2002	960 964-965 974 978-979 984 152 201 498 983 2 20 23 26 28-29 31 37 39 44 57 74 78-79 111 118 122-123 126 129 143 145 148 155 168-169 175-176 178 181 185-186 191 200-202 208 212 214-215 220 222 224 228 230- 231 235 237 239 248-249 252 255- 260 262 266-269 272 280 284 286 289 313 323 326 329 346 358 361 379 396 400 412 422-423 428 437 439 443 445 450 452 457 461 467- 468 479-480 484 487 490 498 500 504-505 523 526 533 541-542 547 561-562 571 574-575 580 605 635 637 640 647 653 655 678 680 711 733 746 761 764 766 771 776 795 865 885 887 900-901 905 907 910 917 924 930 932 941-942 951 958 960 962 967 974-975 979 982-983 989 993 999 1003-1004 23 31 53 87 107 123 160 175 185 197 202 207 215 222 237 252 256- 258 274 284 289 326 358 396 400 437 445 452 462 464 467 487 500 504 526 575 583 590 605 630 653 655 703 733 757 764 795 865 884- 885 900 905 919 924 974-975 982

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY NAME	SEQ ID NOS: OF NUCLEOTIDE(S)
	University	TVILLE	379 764 910 942 951
lung, fibroblast	Strategene	LFB001	13-14 26 78 84 91 98 114 122 148 176 197 204 222 246 251 266 379 387 431 437 441 464 479 484 533 553 571 583 619 645-646 711 739 752 910 926 950 965 978 984
lung tumor	Invitrogen	LGT002	13-14 19 31-32 34-39 43 48 64 67 74 76 87 93 95-96 101 111-112 116 122-123 134 138 142 144-145 147- 148 151-152 160 172 178-179 181- 183 187 191-194 197-198 200-202 205 208 210 218 226 228 234 237 246 248 250-252 254-255 257 260- 262 265 268 274 277-279 289 301 320-321 333 336 343 352 355 358 366-368 371 374 379 391-392 397 400-401 406 410 414 423 431 436 440-441 455-456 458 463-464 468 478-480 484 487 498 503-504 511 519 526-527 529 533 541 553 557 561 570-571 575 578 581 583-586 588-589 597 606 616 619 636 638 640 648 650 652 657 680 700 705- 706 708 716 721-722 729 732 739 744-745 752 762 764 782 795 803 812 816-817 838 863 874 877 906 910-911 922 926 941 951 955 957- 958 962-963 968-969 977-978 982- 983 996-997 1007
lymphocytes	ATCC	LPC001	13-14 35 66 79 95 106-107 112 122-123 149 152 178 181 201 205 246 251-252 267 293 299 358 379 384 400-401 409 415 418 439 443- 444 451 456 458 479 484 487 513 533 568 572 575 583 614 619 686 706 721 730-731 739 747 764 789 905 910 941-942 950 965 978-979 1007
leukocyte	GIBCO	LUCOOI	13-14 19 23 30-32 36 39 45 48-49 60-61 63 67 73-74 78-79 81-82 84 87 91 98-99 107-109 111-112 114 122-123 129 142 144-145 148-150 152 170 176 179 181 183 187-188 194 198 201-208 212-213 215 222 228 235 237 241-242 244-246 249- 251 254-257 263 267 278-280 282- 284 286 289-290 295 302 308-309 313 317 333 337 343 346 356-358 371 379 391-392 394 397 400-401 404 406-410 412-415 423-424 429 431 436 439-441 443-445 450 456 458 479-480 484 487-488 495 498- 500 503 505 511-514 519 523 530- 533 539 541 555 559 561 565-566 570 572 577-578 583 590 595 597 617 619 633 635-636 639-640 646 660 670 672 677 680-681 698 703 705 729 732 739-740 743 747 750 763-764 771 782 792-793 803-805 809 819 838 857 866-867 885 888

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ	SEQ ID NOS: OF NUCLEOTIDE(S)
TISSUE ORIGIN	RNA SOURCE	LIBRARY NAME	SEQ ID NOS: OF MUCLEOIIDE(S)
			900 905 910-911 924 926 928 930
			941 948 950-953 955 962-963 965
			977-979 984 987 989 999 1008
leukocyte	Clontech	LUC003	19 26 68 76 96 122 147 152 198
			201 205 208 284 317 354 358 430
			436 440 479 511 533 541 553 561
			583 589 646 698 732 764 766 838
			984
melanoma from cell line ATCC	Clontech	MEL004	8 23 36 69 91 114 122-123 126 148
#CRL 1424			151 181 202 204 227 246 256-257 265 313 379 391 400 417 466 478-
#CKD 1424			479 487 496 519 521 523 561 570
			583 590 669 728 764 784 838 842
			910 941 950 965 970
mammary gland	Invitrogen	MMG001	4 19 23 26 29 34-39 43 45 48 55
1 3			64 66 74 78 87 96-97 114 116 126
			129 136 142 149 151 155-156 160
			164 168 173 175-176 178 180-181
			183 192 197-200 202 204 207-208
			215 222 226-228 230 232 235-238
			242 246 248 250 252-257 261-262
			268 272 274 278 280 301 303 322
			329 335 337 343 363 368-371 374
			379 381 391 397 400-401 417 426 429 431 437 439-441 443 445 449-
			450 455 464 475 478-479 484-485
			487-488 498-499 504 507 512 517
			519 523 526 532-533 553 557 565
			570-571 573 575 577-578 590-591
			606 617 619 636 640 646 648 663
			677-678 680 691 697 702 708 711
			732 744 764 792 803 811-813 817
			875-877 885 887-888 900 902 905
			908 910-911 918 921-922 934 937
			939 941-942 946 951 958 960 965
2 - 3 5		177777000	968 983 989 993 999 1003 1008
induced neuron cells	Strategene	NTD001	39 122 148 152 181 212 246 266
Cerra			313 337 358 379 452 467 479 484 519 553 561 583 621-626 680 872
			881 910 924 941
retinoid acid	Strategene	NTR001	37 148 152 168 541 583
induced neuronal cells	boracegeme	11110011	37 240 132 100 341 303
neuronal cells	Strategene	NTU001	29 37 147 202 221-222 237 246 262
			337 361 391 400 429 439 460 487
			504 526 541 583 772 816 924 945
			965
pituitary gland	Clontech	PIT004	391 396 764
placenta	Clontech	PLA003	123 183 544 803
prostate	Clontech	PRT001	60-61 76 96 122 145-148 153-154
			175 178 183 201 204 226 228 235
			237 241 245 248 250-251 256 262
			265 280 284 324-325 337 397 400
			409 436-437 456 464 478 480 487 489-490 492 508 516-517 524 552
			561 583 605 722 740 747 849 889
			906 924 926 939 958 974 1005
rectum	Invitrogen	REC001	26 29 43 48 70 74 80 108 114 135-
		100001	136 140 168 178-179 208 226 257

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY	SEQ ID NOS: OF NUCLEOTIDE(S)
 		NAME	262 346 348 371 379 411 413 436-
			437 475 479 484 499 504 517 526
	ļ		534 548-549 555 570 577-578 606
			636 697 729 764 778 793 885 900
	1	1	906 908 910 937 941 951 965 989
			999
salivary gland	Clontech	SAL001	7 38 43 74 87 98 112 122 136 142
			148 162 169 181 183-185 207 215
			228 235 250 254-255 265 280 349-
	1		350 394 437 443 464 508 515-516
			519 559 598 614 619 658 666-667
	1		680 724 762-763 771 803 816 842
			930 933-934 953
salivary gland	Clontech	SALS03	48 108 515 617 900
skin	ATCC	SFB001	39
fibroblast		33333	
skin	ATCC	SFB002	222 803
fibroblast	AICC	515002	
	ATCC	SFB003	237
skin fibroblast	AICC	315003	231
	Glont ach	CTMOOT	16 19 29 39 48 56 65 73 96 108
small	Clontech	SINOOL	
intestine			122 136 148 152 155 160 162 165
			168 172 181 191 208 234 244 246
			266 282 296 379 394 431 440 443
			464 479-480 484 519 571 578 583
			617 619 648 662 694 703 752 763
	•		806 838 908 910 926 937 941 966
			972 976
skeletal	Clontech	SKM001	34 112 116 147 149 152 163 167
muscle			373 379 484 515 553 561-562 781
			838 910 941
spinal cord	Clontech	SPC001	19 22 29 31 55 58 70-71 78 122
_			134 145 148 150 152 159-160 163
			166 171 175-176 183 200-201 203-
			204 220 222 224 235 237 246 248
			250 257 262 266-268 279-280 327-
			328 330 337 343 346 371 379 389
	1		396 416 429-430 437 443 452-453
			456 467 475 479 493-494 498 500
			502 541 544 553 561 583 619 635-
	1	1	
	1	ŀ	636 638 640 680 682 696 764 785
			636 638 640 680 682 696 764 785 900 902 910 941 950 982 994
adult enleen	Clontech	SPI.c01	900 902 910 941 950 982 994
adult spleen	Clontech	SPLC01	900 902 910 941 950 982 994 254 529 701
adult spleen stomach	Clontech Clontech	SPLc01 STO001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178
			900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254
			900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512
			900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921
			900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619
stomach	Clontech	ST0001	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925
stomach	Clontech	STO001 THA002	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925 941 950 967 981 1003
stomach	Clontech	STO001 THA002	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925 941 950 967 981 1003
stomach	Clontech	STO001 THA002	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925 941 950 967 981 1003 29 78 112 122 148 151 160-161 169 176 180-181 183 188 198 201 204- 206 212 250 254 313 374 379 397
stomach	Clontech	STO001 THA002	900 902 910 941 950 982 994 254 529 701 48 53 72 74 122 142 152 161 178 181 200-202 204 208 240 251 254 265 268 309 347 397 410 437 512 539 550 583 616 636 657 659 720 722 921 35 53 78 114 123 156 176 181 228 235 246 252 255-256 265 280 329 331 343 379 437 452 457 467 479 484 496 507 519 553 571 593 619 692 723 754 758 764 853 910 925 941 950 967 981 1003 29 78 112 122 148 151 160-161 169 176 180-181 183 188 198 201 204-

TABLE 1

TISSUE ORIGIN	RNA SOURCE	HYSEQ LIBRARY NAME	SEQ ID NOS: OF NUCLEOTIDE(S)		
			838 910 941-942 944 947 958 969 979 982 989 999 1007		
thymus	Clontech	THMC02	9 19 32 36 63 67 74 78 80 85-86 122-123 138 142 145 147-148 160- 161 169 175-176 181 183-184 187 194 198 202 204 208 211 238 244 246 250 252-254 257 262 265 270- 271 283-285 317 333 349 359-360 379 400-401 406 413 418 429 431 433 436 440-441 473 479 484 487 512-513 517-518 523 525 529 533 535-537 541 544 553 556 561 565 567-570 572-573 578 583 615-619 636 644 660-661 681 683 687 698 732 739 763-764 783 785 789 807- 808 811 816 842 852 864 868-869 900 904 906 910 924 926 930 938 941 965 968 974 979 992 1006-1007		
thyroid gland	Clontech	THROO1	5 10 13-14 19 23 35 37 39 47 59-61 64 74 79 87 100 110 112 117 122-123 133 141-142 145 148 152 156 160 168 181 187 199-202 204-205 207-208 210 220 224-225 228 234-235 237 246-247 251-252 254-256 262 265 267-268 280-281 284 286 301 308 325 332-333 335 337 343 346 363 371 374 378-379 383 394 396-397 400 420 429 431-432 436 445 452 456 464 467-468 474 479-480 484 487 492 499 507 519 522 533 537 550 553 559 561 569 583 619 638 650 653 655 672 678 680 692 705 719 727 748 764 766-767 769 792 797 816 821 854 906 910-911 921 924 926 928 941 946 951 958 960-961 967 971 974-975 978 984 989 999		
trachea	Clontech	TRC001	43 48 108 112 142 148 168 204 208 212 221-222 254 265 282 286 317 371 382 425 440 501 553 565 910		
uterus	Clontech	UTR001	1 37 39 62 145 148 163 183 188 200 257 265 268 346 372 405 408 420 431 520 538 561-562 571 640 680 711 842 850-851 885 910 957		

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	%
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
1	AF208846	Homo sapiens	BM-004	172	43
2	Y53871	Homo sapiens	A human brain-	574	99
			derived signalling		
			factor polypeptide.		
3	AE003620	Drosophila	CG8486 gene product	112	33
		melanogaster			
4	AF193807	Homo sapiens	Rh type B glycoprotein	1204	96
5	Y87156	Homo sapiens	Human secreted	89	46
			protein sequence SEQ ID NO:195.		
6	Y71062	Homo sapiens	Human membrane	135	30
			transport protein, MTRP-7.		
7	AB047936	Macaca	hypothetical	81	38
		fascicularis	protein		
8	Y36156	Homo sapiens	Human secreted	158	68
			protein #28.		
9	AB040964	Homo sapiens	KIAA1531 protein	495	100
10	U29725	Homo sapiens	BMK1 alpha kinase	114	35
11	X00822	Gallus gallus	collagen type III	54	52
12	Y27868	Homo sapiens	Human secreted	119	43
			protein encoded by		
			gene No. 107.		
13	W74813	Homo sapiens	Human secreted	722	92
			protein encoded by		
			gene 85 clone		
14	W74813	Homo sapiens	HSDFV29. Human secreted	722	92
14	W/4813	nomo saprens	protein encoded by	122	1 32
			gene 85 clone		
			HSDFV29.		
15	AF119851	Homo sapiens	PRO1722	333	70
16	AF264750	Homo sapiens	ALR-like protein	133	100
17	X91014	Mus musculus	alpha 1 type XI	131	72
			collagen		
18	AF090930	Homo sapiens	PRO0478	109	90
19	Y86456	Homo sapiens	Human gene 46-	618	95
			encoded protein		
			fragment, SEQ ID		
			NO:371.		
20	AF084535	Homo sapiens	laforin	1809	100
21	Y27585	Homo sapiens	Human secreted	587	98
			protein encoded by		1
		1	gene No. 19.	1214	37
22	268748	Caenorhabditi	Similairity to	214	37
		s elegans	Yeast hypothetical		1
			protein YEH4	1	
			(SW:YEH4_YEAST)~cDN A EST yk87c11.3		
			comes from this		
			gene~cDNA EST		
			yk87c11.5 comes		
		<u> </u>	yko/cii.5 comes	<u> </u>	<u> </u>

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	%
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
			from this gene~cDNA		
			EST yk497d5.3 comes		
			from this gene~cDNA		
	•		EST yk186a5.5 comes		
			from this gene~cDNA		
	-		EST yk243b10.5		
			comes from this		
			gene~cDNA EST		
			yk497d5.5 comes		
			from this gene		
23	D86973	Homo sapiens	similar to Yeast	12053	100
-			translation		
			activator GCN1		
			(P1:A48126)		
24	Y09945	Rattus	putative integral	458	50
		norvegicus	membrane transport	130	
		1.01.031040	protein		
25	U25739	Mus musculus	YSPL-1 form 1	719	77
26	AK024427	Homo sapiens	FLJ00016 protein	668	100
27	AP001707	Homo sapiens	human gene for	603	100
21	AFOULTOT	nomo saprens	claudin-8,	003	100
			Accession No.		
			AJ250711		
28	U16030	Brugia malayi	cuticular collagen	78	37
-			Bmcol-2		37
29	G02479	Homo sapiens	Human secreted	442	100
			protein, SEQ ID NO: 6560.		
30	Y13375	Homo sapiens	Amino acid sequence	1806	99
			of protein PRO262.	1	
31	AF077226	Homo sapiens	copine III	1757	65
32	W75198	Homo sapiens	Human secreted	208	100
		_	protein encoded by		
			gene 3 clone		
			HCEDO84.		
33	AF151978	Homo sapiens	amino acid	3436	100
		1	transporter B0+		
34	Y66735	Homo sapiens	Membrane-bound	1006	100
			protein PRO1153.		
35	AC003093	Homo sapiens	OXYSTEROL-BINDING	764	60
20	110000000	lacino papacino	PROTEIN; 45%	1,01	
			similarity to		
			P22059		
			(PID:g129308)		
36	AF286861	Fasciola	tegumental antigen-	79	30
-3	111 200001	hepatica	like protein	'	
37	AF201945	Homo sapiens	HNOEL-iso	2152	100
38	AF258465		OTRPC4	1668	
		Homo sapiens	.		99
39	AF173003	Homo sapiens	apoptosis regulator	2421	100
40	Y53023	Homo sapiens	Human secreted	128	41
			protein clone		
			qf662_3 protein		
		J	sequence SEQ ID	J	

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
NOCHEOTIDE			NO:52.	BCORB	
41	M25750	Oryctolagus cuniculus	sarcolumenin precursor	2307	97
42	G03797	Homo sapiens	Human secreted protein, SEQ ID NO: 7878.	186	75
43	X57805	Homo sapiens	immunoglobulin lambda light chain	1102	91
44	AE003689	Drosophila melanogaster	CG4596 gene product	419	44
45	Y50934	Homo sapiens	Human fetal brain cDNA clone vc30_1 derived protein #1.	644	100
46	Y19562	Homo sapiens	Amino acid sequence of a human secreted protein.	80	45
47	AF016272	Homo sapiens	Ksp-cadherin	4263	99
48	R13111	Homo sapiens	1B1 IgG aberrant light chain with duplicated variable region.	1000	92
49	AK001636	Homo sapiens	unnamed protein product	1630	97
50	Y65155	Homo sapiens	Human 5' EST related polypeptide SEQ ID NO:1316.	78	34
51	G00471	Homo sapiens	Human secreted protein, SEQ ID NO: 4552.	281	91
52	AJ272050	Homo sapiens	transcription initiation factor IA protein	165	68
53	Y42388	Homo sapiens	Amino acid sequence of pt127_1.	668	73
54	AF193807	Homo sapiens	Rh type B glycoprotein	248	97
55	AF132611	Homo sapiens	monocarboxylate transporter MCT3	139	37
56	U43940	Rattus norvegicus	focal adhesion kinase	141	84
57	L17318	Rattus norvegicus	proline-rich proteoglycan	124	37
58	G02832	Homo sapiens	Human secreted protein, SEQ ID NO: 6913.	132	48
59	G00357	Homo sapiens	Human secreted protein, SEQ ID NO: 4438.	95	64
60	¥12723	Homo sapiens	Human 5' EST secreted protein SEQ ID NO:313.	91	50
61	Y19450	Homo sapiens	Amino acid sequence of a human secreted	406	100

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	왕
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
	755555		protein.		
62	AF156549	Mus musculus	putative E1-E2 ATPase	876	65
63	AL356276	Homo sapiens	bA367J7.5 (novel	655	84
			Immunoglobulin	Ì	
			domain containing		
			protein)		ļ
64	AL133105	Homo sapiens	hypothetical protein	1783	99
65	U32189	Oryctolagus	histidine-rich	73	40
		cuniculus	glycoprotein		
			precursor		
66	Y91433	Homo sapiens	Human secreted	758	98
			protein sequence		
			encoded by gene 33		
			SEQ ID NO:154.	<u> </u>	
67	W75198	Homo sapiens	Human secreted	208	100
		1	protein encoded by		
			gene 3 clone		
			HCEDO84.		
68	AF020651	Homo sapiens	T cell receptor	742	93
			alpha chain		
	7.57.1000.0	-:	variable region		
69	AF118086	Homo sapiens	PRO1992	158	61
70	X52454	Drosophila melanogaster	rho	224	36
71	W40353	Homo sapiens	Human unspecified	146	67
			protein from		1
			US5702907.		
72	Y66690	Homo sapiens	Membrane-bound	971	98
			protein PRO813.		
73	AJ002744	Homo sapiens	UDP-	1518	98
			GalNAc:polypeptide		
			N-		
			acetylgalactosaminy		
74	AC024792	Caenorhabditi	ltransferase 7	423	
/ 4	AC024732	s elegans	contains similarity to TR:P78316	423	36
75	AB016088	Homo sapiens	RNA binding protein	109	32
76	Y94953	Homo sapiens	Human secreted	2484	100
70	154553	nomo saprens	protein clone	2404	100
			fy356 14 protein		
			sequence SEQ ID		
			NO:112.		
77	AF107406	Homo sapiens	GW128	74	51
78	Y13401	Homo sapiens	Amino acid sequence	1681	96
			of protein PRO339.		:
79	Y94290	Homo sapiens	Human myosin heavy	1819	99
9.0	7 500 77 04	Lione	chain homologue.	- <u></u>	
80	AF007194	Homo sapiens	mucin	4875	100
81	AF229179	Homo sapiens	kidney-specific	949	99
			membrane protein		1
		<u> </u>	NX-17	L	L

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
82	AL356173	Neurospora crassa	hypothetical protein	83	29
83	G00437	Homo sapiens	Human secreted protein, SEQ ID NO: 4518.	87	69
84	K03036	Mus musculus	alpha-1 type I procollagen	114	38
85	AF233261	Homo sapiens	otoraplin	676	100
86	AF073519	Homo sapiens	small EDRK-rich factor 1, long isoform	100	45
87	AC021640	Arabidopsis thaliana	putative phosphatidate phosphohydrolase	387	43
88	AB040812	Homo sapiens	protein kinase PAK5	1159	100
89	AL365409	Homo sapiens	similar to (NP_034322.1) sex- determination protein homolog Femla	694	100
90	U81035	Rattus norvegicus	ankyrin binding cell adhesion molecule neurofascin	189	63
91	W88684	Homo sapiens	Secreted protein encoded by gene 151 clone HNHED86.	134	65
92	Y66734	Homo sapiens	Membrane-bound protein PRO1097.	297	70
93	AB031051	Homo sapiens	organic anion transporter OATP-E	283	40
94	B08976	Homo sapiens	Human secreted protein sequence encoded by gene 28 SEQ ID NO:133.	71	27
95	U83115	Homo sapiens	non-lens beta gamma-crystallin like protein	245	97
96	AF156551	Mus musculus	putative E1-E2 ATPase	3779	86
97	AF062476	Mus musculus	retinoic acid- responsive protein; STRA6	1091	74
98	Y87072	Homo sapiens	Human secreted protein sequence SEQ ID NO:111.	490	100
99	AF116652	Homo sapiens	PRO0813	1015	99
100	AF159567	Homo sapiens	C2H2 (Kruppel-type) zinc finger protein	2176	100
101	D25328	Homo sapiens	platelet-type phosphofructokinase	109	95
102	AB018563	Homo sapiens	TML1	98	68
103	X83107	Homo sapiens	bmx	232	85

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
104	U49973	Homo sapiens	ORF1; MER37;	131	43
101	045575	nomo bapiens	putative	131	1 = 3
			transposase similar		
			to pogo element		
105	Y86472	Homo sapiens	Human gene 52-	150	54
100	1804/2	nomo saptens	encoded protein	1 130	24
		5	fragment, SEQ ID		
			NO:387.		
106	AF020276	Homo sapiens	spinocerebellar	96	37
100	141 020270	nomo saprema	ataxia 7	30	37
107	W57901	Homo sapiens	Protein of clone	1499	96
107	W3 / 301	nomo saprens	CT748 2.	1433	96
108	R13111	Homo sapiens	1B1 IgG aberrant	1210	84
108	KISILI	HOMO Saptems	light chain with	1210	04
			duplicated variable		
			region.		
109	W50192	Homo sapiens	Amino acid sequence	95	32
103	W50192	nomo saprens	of salivary protein	35	32
			CON-1.		
110	AB046634	Macaca	hypothetical	282	75
110	ABU46634	Macaca fascicularis	protein	282	/5
111	AF242432	Mus musculus	neuronal apoptosis	486	30
TTT	AF242432	Mus muscurus		486	29
			inhibitory protein		
110	AB000280	Rattus	peptide/histidine	2400	-
112	AB000280		1	2490	88
113	AF182443	norvegicus Rattus	transporter	597	99
112	AF182443		F-box protein FBL2	597	99
114	AJ245874	norvegicus	7 77 /	7.0.10	
774	AU245874	Homo sapiens	putative ATG/GTP	1242	100
115	AF179828	Saimiri	binding protein	1	-
112	AF1/9828	sciureus	olfactory receptor	444	66
116	Y66735		Membrane-bound	1006	100
110	166/35	Homo sapiens	protein PRO1153.	1006	100
117	Y94344	Homo sapiens	Human cell surface	892	90
11/	194344	HOMO Sapiens	receptor protein	892	90
			#11.		
118	AJ238706	Drosophila	monocarboxylate	226	31
110	AU238706	ı -	transporter 1	226	37
		melanogaster	homologue		
110	77700700	B	<u> </u>	27.0	
119	AF180728	Drosophila	sulfate transporter	312	4.5
100	1 77004000	melanogaster	7 1	 	
120	AE004890	Pseudomonas	L-lactate permease	534	89
101	1	aeruginosa	ļ		
121	X91837	Saccharomyces	cell division cycle	435	98
100	1 1100555	cerevisiae	protein CDC55		
122	U93565	Homo sapiens	putative p150	1911	90
123	AJ000332	Homo sapiens	Glucosidase II	5043	99
124	AF204674	Homo sapiens	muscle disease-	377	72
			related protein		
125	S58722	Homo sapiens	X-linked	196	68
			retinopathy protein		
	1	I	{C-terminal, clone		1

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	્રે .
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
NOCHBOTTER			XEH.8c		
126	\$58722	Homo sapiens	X-linked retinopathy protein {C-terminal, clone XEH.8c}	196	68
127	J03848	Mesocricetus auratus	metallothionein II	147	51
128	G02994	Homo sapiens	Human secreted protein, SEQ ID NO: 7075.	93	64
129	AF116238	Homo sapiens	pseudouridine synthase 1	1927	99
130	G03411	Homo sapiens	Human secreted protein, SEQ ID NO: 7492.	183	65
131	AF222861	Sus scrofa	type X collagen	90	34
132	G03628	Homo sapiens	Human secreted protein, SEQ ID NO: 7709.	60	66
133	Y10529	Homo sapiens	olfactory receptor	766	61
134	AF164612	Homo sapiens	Gag protein	125	43
135	Y12713	Mus musculus	Pro-Pol-dUTPase polyprotein	181	47
136	X57816	Homo sapiens	immunoglobulin lambda light chain	550	57
137	U07808	Mus musculus	metallothionein IV	55	37
138	AB031227	Pisum sativum	PsAD1	68	50
139	AB035520	Oryctolagus cuniculus	parchorin	1324	57
140	AB007891	Homo sapiens	KIAA0431	117	46
141	Y00278	Homo sapiens	Human secreted protein encoded by gene 21.	234	92
142	Y68810	Homo sapiens	A rat heavy chain region and a human hinge region.	1124	92
143	M58526	Homo sapiens	alpha-5 type IV collagen	4597	97
144	AF119851	Homo sapiens	PRO1722	192	66
145	X84908	Homo sapiens	phosphorylase kinase	3798	97
146	Y76155	Homo sapiens	Human secreted protein encoded by gene 32.	81	52
147	U13766	Murine . leukemia virus	gag-pol polyprotein	735	36
148	AF034198	Homo sapiens	IGSF1	7154	100
149	Y94343	Homo sapiens	Human cell surface receptor protein #10.	1331	100
150	Y87211	Homo sapiens	Human secreted	759	97
			I		

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	9
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
			protein sequence SEQ ID NO:250.		
151	AJ252258	human	glycoprotein G-2	115	30
	1333333	herpesvirus 2	grycoprocesm o z		
152	V00662	Homo sapiens	URF 1 (NADH	1283	85
			dehydrogenase		
			subunit)		
153	G02872	Homo sapiens	Human secreted	142	61
			protein, SEQ ID NO:		
154	722706	D-1	6953.		
155	A23786 Z34465	Beta vulgaris Zea mays	chitinase 1 extensin-like	138 97	41
133	234465	Zea mays	protein	97	36
156	X79389	Homo sapiens	glutathione	721	66
	11,3303	nomo baprens	transferase Tl	1 / 2 1	00
157	M22333	Homo sapiens	unknown protein	106	46
158	AL118502	Homo sapiens	bA371L19.1 (novel	2471	100
			protein)		
159	AJ012582	Homo sapiens	hyperpolarization-	3076	100
			activated cation		
			channel HCN2		
160	D26351	Homo sapiens	human type 3	8901	99
			inositol 1,4,5- trisphosphate		!
			receptor		
161	AF067656	Homo sapiens	ZW10 interactor	951	97
			Zwint	332	"
162	AE003461	Drosophila	CG11300 gene	76	29
		melanogaster	product		
163	Y48518	Homo sapiens	Human breast	355	100
			tumour-associated		ĺ
			protein 63.		
164	G00517	Homo sapiens	Human secreted	83	34
			protein, SEQ ID NO: 4598.		
165	G03786	Homo sapiens	Human secreted	251	53
		nomo bapacino	protein, SEQ ID NO:	232	
			7867.		
166	Y00765	Homo sapiens	Prion protein CJAS.	63	37
167	Y21050	Homo sapiens	Human glial	206	71
			fibrillary acidic		
			protein GFAP mutant		
			fragment 59.		
168	X74929	Homo sapiens	Keratin 8	1462	95
169	U29488	Caenorhabditi	similar to DNAJ	555	29
170	L27428	s elegans Homo sapiens	protein reverse	145	45
_, _	12/420	10000 saptens	reverse transcriptase	T#2	45
171	W19932	Homo sapiens	Alzheimer's disease	362	100
		Suprems	protein encoded by	302	1.00
	•		DNA from plasmid		
			pGCS55.		
172	AF178983	Homo sapiens	Ras-associated	497	100

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
			protein Rap1		
173	U70136	Homo sapiens	megakaryocyte stimulating factor; MSF	206	28
174	G00352	Homo sapiens	Human secreted protein, SEQ ID NO: 4433.	109	64
175	U28143	Gallus gallus	synemin	1014	39
176	Y13401	Homo sapiens	Amino acid sequence of protein PRO339.	1978	96
177	AJ243396	Homo sapiens	voltage-gated sodium channel beta-3 subunit	947	99
178	M77812	Oryctolagus cuniculus	myosin heavy chain	4079	98
179	AF200344	Homo sapiens	aspartyl protease 3	956	91
180	AF200815	Homo sapiens	FUSED serine/threonine kinase	1597	99
181	G03786	Homo sapiens	Human secreted protein, SEQ ID NO: 7867.	147	83
182	Y00313	Homo sapiens	Human secreted protein encoded by gene 56.	56	29
183	X00699	Homo sapiens	precursor	583	66
184	AF269289	Homo sapiens	unknown	81	32
185	G03797	Homo sapiens	Human secreted protein, SEQ ID NO: 7878.	176	66
186	Y20298	Homo sapiens	Human apolipoprotein E mutant protein fragment 11.	110	34
187	AF161437	Homo sapiens	HSPC319	867	99
188	Y19684	Homo sapiens	SEQ ID NO 402 from W09922243.	124	47
189	¥74050	Homo sapiens	Human prostate tumor EST fragment derived protein #237.	78	42
190	Y08986 ,	Brassica napus	oleosin-like protein	106	36
191	AF119851	Homo sapiens	PRO1722	173	66
192	AF116712	Homo sapiens	PRO2738	166	50
193	AF186084	Homo sapiens	epidermal growth factor repeat containing protein	2022	85
194	M59819	Homo sapiens	granulocyte colony- stimulating factor receptor	4232	100
195	Y86228	Homo sapiens	Human secreted protein HFXJX44,	250	100

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	ે
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			SEQ ID NO:143.		
196	Y45382	Homo sapiens	Human secreted	181	63
			protein fragment	,	
			encoded from gene		1
408			28.	ļ	
197	X94991	Homo sapiens	zyxin	566	41
198	M17236	Homo sapiens	MHC HLA-DQ alpha precursor	896	84
199	AC004659	Homo sapiens	BC62940_2	805	53
200	X14420	Homo sapiens	prepro-alpha-1 type 3 collagen	5521	99
201	AF180473	Homo sapiens	Not2p	1628	98
202	X85237	Homo sapiens	human splicing	1145	100
			factor		
203	AL390114	Leishmania	extremely	309	58
İ		major	cysteine/valine	į	
			rich protein		
204	D42138	Homo sapiens	PIG-B	1479	98
205	Y00062	Homo sapiens	precursor	3334	98
			polypeptide (AA -23 to 1120)		
206	W93946	Homo sapiens	Human regulatory	1011	100
			molecule HRM-2		
	<u></u>		protein.		
207	AB017563	Homo sapiens	IGSF4	2062	99
208	X54637	Homo sapiens	protein tyrosine kinase	5694	98
209	AF255910	Homo sapiens	vascular	1508	98
			endothelial		
			junction-associated molecule		
210	AF061324	Homo sapiens	sulfonylurea	7545	97
		1	receptor 2A		
211	U93568	Homo sapiens	p40	197	50
212	AF250842	Drosophila	multiple asters	506	32
		melanogaster			
213	X81479	Homo sapiens	EMR1	4469	99
214	X77748	Homo sapiens	metabotropic	4471	99
			glutamate receptor		
			type 3 (mGluR3)		
215	M60396	Homo sapiens	transcobalamin II	2218	99
216	W48351	Homo sapiens	Human breast cancer	170	71
			related protein		
	152.50.62		BCRB2.		
217	Y36203	Homo sapiens	Human secreted protein #75.	156	73
218	AF119851	Homo sapiens	PRO1722	144	63
219	AJ246002	Mus musculus	spastin protein	143	100
			orthologue		
220	D49958	Homo sapiens	membrane glycoprotein M6	616	57
221	X83573	Homo sapiens	ARSE	2114	93
	<u> </u>				

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	ક
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	,
222	AF126062	Homo sapiens	Arf-like 2 binding	508	84
		1	protein BART1		
223	L22695	Canine oral	5' end derived by	83	51
225		papillomaviru	splicing; putative		
		s	paragraph paragraph		
224	R95913	Homo sapiens	Neural thread	262	64
224	ROSOIS	nomo suprems	protein.	202	"
225	AP001306	Arabidopsis	contains similarity	79	34
225	APOULSUS	thaliana	to cell wall-plasma	1 ' 3	34
•	1	Charrana	membrane linker		
					1
			protein-gene_id:MKA		
		ļ	23.3		
226	G01984	Homo sapiens	Human secreted	252	64
	İ		protein, SEQ ID NO:		
			6065.		
227	X04614	human	IE110	83	35
		herpesvirus 1			
228	AF151877	Homo sapiens	CGI-119 protein	1203	94
229	AF181467	Homo sapiens	protein Z-dependent	1483	88
			protease inhibitor		
			precursor		
230	Z81326	Homo sapiens	neuroserpin	1763	99
231	AF111173	Homo sapiens	sodium/hydrogen	3512	99
			exchanger isoform 5		
232	X67055	Homo sapiens	inter-alpha-trypsin	4429	98
233	110,033	nome papers	inhibitor heavy		
			chain H3		
233	AB004064	Homo sapiens	tomoregulin	1783	98
234	AL096772	Homo sapiens	dJ365012.1	5465	98
234	AL096772	HOMO SAPIEMS	(KIAA0758 protein)	3403	1 30
235	X83378	Homo sapiens	putative chloride	1620	99
235	A83378	nomo saprens	channel	1620	99
	77042644	 	receptor protein	5127	97
236	AF043644	Homo sapiens	tyrosine	5127	" '
			phosphatase		
			nucleotide binding	1372	100
237	AF208536	Homo sapiens		13/2	100
			protein; NBP		
238	AC005625	Homo sapiens	R27328_1	2435	93
239	X55687	Lycopersicon	extensin (class II)	58	50
		esculentum			
240	M23315	Sesbania	nodulin	61	36
		rostrata	l		
241	AF102851	Homo sapiens	dolichyl-P-	1881	99
			Glc:Man9GlcNAc2-PP-		
			dolichyl		
			glucosyltransferase		
242	G03793	Homo sapiens	Human secreted	202	67
			protein, SEQ ID NO:		
			7874.		
243	G03258	Homo sapiens	Human secreted	203	69
473	303236	TOMO Saprens	protein, SEQ ID NO:	1203	"
			7339.		
1244	AE049774	Home carriers	anti-HER3 scFv	903	81
244	AF048774	Homo sapiens	and neks SCFV	303	01

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
245	AF102851	Homo sapiens	dolichyl-P-	1867	98
			Glc:Man9GlcNAc2-PP-		
			dolichyl		
			glucosyltransferase		
246	L00352	Homo sapiens	low density	3980	100
			lipoprotein		
			receptor		
247	Y79510	Homo sapiens	Human carbohydrate-	1394	100
			associated protein		
			CRBAP-6.		
248	AF202636	Homo sapiens	angiopoietin-like	2164	100
			protein PP1158		
249	X66533	Homo sapiens	guanylate cyclase	1641	97
250	M20504	Homo sapiens	MHC HLA-DR-beta-2	750	70
0.51	1 77 5 77 6 7		precursor	1050	
251	AF157326	Homo sapiens	TIP120 protein	4278	99
252	M25865	Homo sapiens	von Willebrand	10841	95
<u> </u>	7 500 5 60 5	<u> </u>	factor		
253	AC005625	Homo sapiens	R27328_1	2435	93
254	A21385	synthetic construct	heavy chain	1786	94
255	AF182414		antibody 3D6	310	1
256	Y54041	Homo sapiens		1267	48
256	154041	Homo sapiens	Protein encoded by	1267	84
			a gene reduced in metastatic melanoma		
	i		cells (grmm-1).		
257	AJ011415	Homo sapiens	plexin-B1/SEP	1580	60
237	MOOTIFIE	nomo saprens	receptor	1380	80
258	W55030	Homo sapiens	G-protein coupled	1493	100
250		nomo saprens	receptor, long	1 1 1 2 3	100
			form.		
259	AF227747	Homo sapiens	voltage-dependent	6158	100
			calcium channel		====
			alpha 1G subunit		
			isoform bc		
260	AF111173	Homo sapiens	sodium/hydrogen	3512	99
		_	exchanger isoform 5	İ	
261	G01984	Homo sapiens	Human secreted	175	70
		_	protein, SEQ ID NO:		
			6065.		
262	Y00815	Homo sapiens	put. LAR preprotein	5648	100
			(AA -16 to 1881)	ł	
263	234979	Homo sapiens	Human FIZZ3	582	100
			(inhibitor of		
			neurotrophin		
			action) cDNA.		1
264	AF119851	Homo sapiens	PRO1722	189	73
265	AL049798	Homo sapiens	dJ797M17.1	1007	99
		_	(Dermatopontin)		
266	AL035684	Homo sapiens	dJ1114A1.1	1978	99
			(KIAA0611 (putative		1
			E1-E2 ATPase)		
			protein)]	I

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER	,		WATERMAN SCORE	IDENTITY
267	U49055	Rattus norvegicus	rA8	4382	87
268	X15332	Homo sapiens	alpha-1 (III) collagen	4170	99
269	Z98884	Homo sapiens	dJ467L1.1 (KIAA0833)	2010	100
270	AF085244	Homo sapiens	C2H2 type Kruppel- like zinc finger protein splice variant b	7331	98
271	Y00319	Homo sapiens	Human secreted protein encoded by gene 63.	214	82
272	X04434	Homo sapiens	IGF-I receptor	5832	99
273	AC005626	Homo sapiens	R29124_1	1129	89
274	X52046	Mus musculus	type III collagen	819	37
275	M22207	Tripneustes gratilla	217g protein	168	51
276	M32317	Homo sapiens	HLA protein allele B7	1536	84
277	L05485	Homo sapiens	surfactant protein	1693	87
278	W88504	Homo sapiens	Human epidermoid carcinoma clone HP10428-encoded membrane protein.	1187	100
279	AF078850	Homo sapiens	steroid dehydrogenase homolog	794	عست
280	X83378	Homo sapiens	putative chloride channel	1620	99
281	AL035701	Homo sapiens	dJ8B1.3 (similar to PLASMA-CELL MEMBRANE GLYCOPROTEIN PC-1)	2412	99
282	Y87068	Homo sapiens	Human secreted protein sequence SEQ ID NO:107.	528	100
283	L40806	Neurospora crassa	Restriction enzyme inactivation of met-10 complementation in this region. Sequence similarity to S. cerevisiae chromosome VIII cosmid 9205, accession no. U10556 CDS residues 22627-24126	536	35
284	W88552	Homo sapiens	Secreted protein encoded by gene 19 clone HSAVU34.	3078	99

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
285	G03790	Homo sapiens	Human secreted protein, SEQ ID NO: 7871.	108	50
286	X68060	Homo sapiens	DNA topoisomerase	8296	99
287	G00352	Homo sapiens	Human secreted protein, SEQ ID NO: 4433.	114	41
288	AC004602	Homo sapiens	F23487_2	202	49
289	AF196329	Homo sapiens	triggering receptor expressed on monocytes 1	1211	99
290	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	202	62
291	G03043	Homo sapiens	Human secreted protein, SEQ ID NO: 7124.	93	62
292	Y12550	Homo sapiens	Human 5' EST secreted protein SEQ ID NO: 215 from WO 9906553.	141	100
293	D43756	Canis familiaris	fibrinogen A-alpha- chain	102	33
294	U38545	Homo sapiens	phospholipase D1	5681	99
295	W42076	Homo sapiens	The amino acid sequence of the O276 16 protein.	236	100
296	AF090930	Homo sapiens	PRO0478	128	60
297	Y64747	Homo sapiens	Human 5' EST related polypeptide SEQ ID NO:908.	471	98
298	G01234	Homo sapiens	Human secreted protein, SEQ ID NO: 5315.	280	71
299	G02514	Homo sapiens	Human secreted protein, SEQ ID NO: 6595.	94	76
300	G02493	Homo sapiens	Human secreted protein, SEQ ID NO: 6574.	112	46
301	238061	Saccharomyces cerevisiae	mal5, sta1, len: 1367, CAI: 0.3, AMYH_YEAST P08640 GLUCOAMYLASE S1 (EC 3.2.1.3)	340	27
302	Y59672	Homo sapiens	Secreted protein 108-006-5-0-E6-FL.	530	78
303	Y95018	Homo sapiens	Human secreted protein vp19_1, SEQ ID NO:76.	76	35
304	W34623	Homo sapiens	Human C3 protein mutant FT-1.	117	46

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	ો
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE	V05000		**	SCORE	
305	Y87292	Homo sapiens	Human signal peptide containing	81	50
			protein HSPP-69 SEO		
			ID NO:69.		}
306	AF210651	Homo sapiens	NAG18	135	60
307	Y14482	Homo sapiens	Fragment of human	212	58
		_	secreted protein		
			encoded by gene 17.		
308	Y76325	Homo sapiens	Fragment of human	343	93
			secreted protein		
			encoded by gene 35.	l	
309	Y36156	Homo sapiens	Human secreted	203	75
			protein #28.		
310	AF090931	Homo sapiens	PRO0483	76	50
311	AC004943	Homo sapiens	alpha-fetoprotein	351	85
			enhancer-binding		
			protein; 99% identical to A41948		
			(PID:g283975)		
312	G02558	Homo sapiens	Human secreted	144	52
312	002338	nomo saprens	protein, SEQ ID NO:	144	32
			6639.		
313	AK000128	Homo sapiens	unnamed protein	1338	100
			product		
314	G03786	Homo sapiens	Human secreted	164	83
			protein, SEQ ID NO:		
			7867.		İ
315	AF090942	Homo sapiens	PRO0657	253	68
316	AF116712	Homo sapiens	PRO2738	181	52
317	AF043726	Mus musculus	PHD-finger protein	1605	64
318	Y99368	Homo sapiens	Human PRO1326	145	51
			(UNQ686) amino acid		
			sequence SEQ ID NO:100.		
319	AF065314	Homo sapiens	cone photoreceptor	292	98
349	Aroossia	nomo saprens	cGMP-gated channel	252	30
			alpha subunit		
320	AF003389	Caenorhabditi	contains similarity	162	28
		s elegans	to N-chimaerins		
321	Y66755	Homo sapiens	Membrane-bound	993	100
			protein PRO1185.		
322	AF109906	Mus musculus	RD	118	69
323	AF199323	Rattus	RIM2-2A	364	85
	<u> </u>	norvegicus			
324	G02538	Homo sapiens	Human secreted	104	65
			protein, SEQ ID NO:		
			6619.		
325	G02872	Homo sapiens	Human secreted	138	65
			protein, SEQ ID NO:		
332	1	 	6953.	<u> </u>	
326	Y41266	Homo sapiens	Human T139 protein.	591	100
327	G02920	Homo sapiens	Human secreted	103	67
	<u> </u>	1	protein, SEQ ID NO:		

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	%
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	100000
	 		7001.		
328	G00636	Homo sapiens	Human secreted	80	36
323	000000	nomo saprens	protein, SEQ ID NO:		36
	·		4717.		
329	U37769	Oryctolagus	protein phosphatase	556	100
349	03//09	cuniculus		556	88
		cuniculus	2A0 B' regulatory		
İ			subunit alpha	•	
			isoform		
330	AE001424	Plasmodium	RESA-H3 antigen	208	21
		falciparum			
331	AF090930	Homo sapiens	PRO0478	156	82
332	AF161356	Homo sapiens	HSPC093	169	64
333	G04055	Homo sapiens	Human secreted	425	100
			protein, SEQ ID NO:		
			8136.		
334	D79985	Homo sapiens	putative	371	86
			hydrophobic domain		
	i		in the central		
			region.	•	
335	Y41401	Homo sapiens	Human secreted	392	100
			protein encoded by		
			gene 94 clone	İ	
1			HLYCH68.		
336	W18651	Homo sapiens	Human	478	88
		and Daptoni	apolipoprotein E	1 - 7 - 0	00
			gene +1 frameshift		
			mutant product.		
337	Y20921	Homo sapiens	Human presenilin II	2126	96
	150522	nomo bapiono	wild type protein	1 2120	36
			fragment 5.	}	
338	AF010144	Homo sapiens	neuronal thread	233	75
330	MIGIOITA	nomo saprens	protein AD7c-NTP	233	/ 5
339	D28500	Homo sapiens	mitochondrial	175	89
333	D28300	HOMO Sapiens	isoleucine tRNA	1/5	89
					1
340	Y13357	77	synthetase		
340	113357	Homo sapiens	Amino acid sequence	148	50
241	77.006677	777	of protein PRO227.	<u> </u>	
341	AL096677	Homo sapiens	dJ322G13.2 (similar	94	50
343	371.00.43	**	to cystatin)	ļ	
342	Y10843	Homo sapiens	Amino acid sequence	186	86
			of a human secreted		
			protein.		
343	X54134	Homo sapiens	protein-tyrosine	3705	100
			phosphatase		
344	Z33908	Mus musculus	inositol 1,4,5-	315	84
			trisphosphate		
			receptor		
345	G00241	Homo sapiens	Human secreted	130	46
			protein, SEQ ID NO:		
			4322.		
346	AF071172	Homo sapiens	HERC2	23705	99
347	AB015346	Homo sapiens	Eps15R	209	95
348	Y48596	Homo sapiens	Human breast	108	34
		Paprens	Transatt Na Cabl	1 100	

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	identity
			tumour-associated protein 57.		
349	G03058	Homo sapiens	Human secreted protein, SEQ ID NO: 7139.	85	66
350	¥73443	Homo sapiens	Human secreted protein clone yb187_1 protein sequence SEQ ID NO:108.	90	36
351	G03793	Homo sapiens	Human secreted protein, SEQ ID NO: 7874.	126	66
352	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	324	73
353	Y64747	Homo sapiens	Human 5' EST related polypeptide SEQ ID NO:908.	527	98
354	AF255342	Homo sapiens	putative pheromone receptor V1RL1 long form	147	59
355	W48351	Homo sapiens	Human breast cancer related protein BCRB2.	85	61
356	G03060	Homo sapiens	Human secreted protein, SEQ ID NO: 7141.	191	72
357	AF124729	Mus musculus	acinusS'	124	31
358	U37352	Homo sapiens	protein phosphatase 2A B'alphal regulatory subunit	1016	95
359	AF280605	Triticum aestivum	omega gliadin storage protein	125	35
360	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	150	81
361	AL035398	Homo sapiens	dJ796I17.2 (CGI-51)	226	64
362	AK000307	Homo sapiens	unnamed protein product	882	97
363	Y41401	Homo sapiens	Human secreted protein encoded by gene 94 clone HLYCH68.	392	100
364	AF288480	Homo sapiens	tubby super-family protein	238	87
365	AL023706	Schizosacchar omyces pombe	possible pre-mRNA processing by similarity to yeast prp39	383	34
366	W48351	Homo sapiens	Human breast cancer related protein BCRB2.	85	61

TABLE 2

SEQ J.D NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	0,0
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
367	S68978	Oryctolagus	interleukin-1	53	58
		cuniculus	receptor antagonist		
		-	intracellular form		
368	AF047602	Equus zebra	luteinizing	68	37
		hartmannae	hormone/chorionic		
			gonadotrophin beta-		
			subunit		
369	AF119851	Homo sapiens	PRO1722	180	75
370	U15195	Homo sapiens	alpha-1 type II	59	43
			collagen		
371	U02082	Homo sapiens	guanine nucleotide	2648	100
			regulatory protein		
372	AF096895	Homo sapiens	chemokine-like	508	100
			factor 1		
373	G03786	Homo sapiens	Human secreted	315	65
			protein, SEQ ID NO:		İ
			7867.		<u></u>
374	AF010144	Homo sapiens	neuronal thread	240	67
			protein AD7c-NTP		<u> </u>
375	U22376	Homo sapiens	alternatively	191	80
			spliced product		
0.00			using exon 13A		
376	U08310	Saimiri	prion protein	245	66
		sciureus			
377	A76867	unidentified	Chimere G.CSF-Gly4-	550	99
			SAH en aval region	1	
378	G00442	77	prepro de SAH		
3/8	G00442	Homo sapiens	Human secreted	94	53
		†	protein, SEQ ID NO: 4523.		
379	AF010144	Homo sapiens	neuronal thread	\	
3,3	ALOIOI44	nomo saprens	protein AD7c-NTP	355	53
380	AB023634	Rattus	Ca/calmodulin-	161	91
300	AD023034	norvegicus	dependent protein	101	91
		noi vegicus	kinase phosphatase		
381	Y99437	Homo sapiens	Human PRO1508	805	100
- 		Marie Bupiens	(UNQ761) amino acid	803	100
			sequence SEQ ID		
			NO:336.		
382	W48351	Homo sapiens	Human breast cancer	139	61
	1		related protein		
			BCRB2.		
383	M58511	Homo sapiens	iron-responsive	286	100
			element-binding		
			protein/iron		
			regulatory protein		
			2	1	
384	Y02671	Homo sapiens	Human secreted	99	71
	1		protein encoded by		
	1		gene 22 clone	1	
			HMSJW18.		
385	AJ012166	Canis	brain-specific	86	38
303	130022200	COLLEGE		100 1	J0

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	ક
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			protein, Bassoon		
386	L07809	Homo sapiens	dynamin	98	31
387	M15530	Homo sapiens	B-cell growth	158	69
			factor		
388	AF090172	Mycoplasma	revertant adhesin-	109	31
		pneumoniae	related protein P30		
389	AJ278964	Homo sapiens	cytosolic beta- glucosidase	165	52
390	AF190642	Homo sapiens	phosphoinositide- specific phospholipase C PLC-epsilon	1095	98
391	X13238	Homo sapiens	cytochrome c oxidase subunit VIc preprotein	379	100
392	AF225417	Homo sapiens	88.8 kDa protein	1634	98
393	Y02693	Homo sapiens	Human secreted protein encoded by gene 44 clone HTDAD22.	278	75
394	AF151037	Homo sapiens	HSPC203	554	100
395	AJ276396	Homo sapiens	matrix extracellular phosphoglycoprotein	465	100
396	X51405	Homo sapiens	pre-pro polypeptide (AA -25 to 451)	2536	100
397	W78128	Homo sapiens	Human secreted protein encoded by gene 3 clone HOSBI96.	564	71
398	¥87346	Homo sapiens	Human signal peptide containing protein HSPP-123 SEQ ID NO:123.	290	90
399	G03564	Homo sapiens	Human secreted protein, SEQ ID NO: 7645.	72	52
400	U89436	Homo sapiens	tyrosyl-tRNA synthetase	2719	100
401	W80993	Homo sapiens	Human RIP- interacting factor RIF.	1724	100
402	Y27907	Homo sapiens	Human secreted protein encoded by gene No. 119.	95	59
403	AB033102	Homo sapiens	KIAA1276 protein	921	100
404	G03797	Homo sapiens	Human secreted protein, SEQ ID NO: 7878.	192	55
405	AF096895	Homo sapiens	chemokine-like factor 1	508	100
406	¥29861	Homo sapiens	Human secreted protein clone	791	98

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	IDENTITY
·			cb98_4.		
407	Y00293	Homo sapiens	Human secreted protein encoded by gene 36.	237	97
408	W40215	Homo sapiens	Human macrophage antigen.	1358	99
409	L36056	Homo sapiens	4E-binding protein 2	639	100
410	AJ130710	Homo sapiens	QA79 membrane protein, allelic variant airm-lb	2473	100
411	AF116661	Homo sapiens	PRO1438	146	57
412	W88761	Homo sapiens	Polypeptide fragment encoded by gene 19.	150	58
413	AK024434	Homo sapiens	FLJ00024 protein	574	97
414	Y10376	Homo sapiens	SIRP-betal	2069	99
415	Y07930	Homo sapiens	Human secreted protein fragment encoded from gene 79.	351	98
416	R99390	Homo sapiens	Human 030 gene (fohy030) product.	804	71
417	AB018253	Rattus norvegicus	voltage-gated ca channel	2419	88
418	AC006017	Homo sapiens	similar to ALR; similar to AAC51735 (PID:g2358287)	2150	97
419	X72925	Homo sapiens	Dsc1b precursor	4390	99
420	AF205940	Homo sapiens	endomucin	1289	100
421	Y27868	Homo sapiens	Human secreted protein encoded by gene No. 107.	134	54
422	W74722	Homo sapiens	Human secreted protein er80_1.	2422	100
423	AF080470	Homo sapiens	pallid	872	100
424	G04072	Homo sapiens	Human secreted protein, SEQ ID NO: 8153.	201	63
425	W90961	Homo sapiens	Human CSGP-1 protein.	869	86
426	M13180	Human herpesvirus 4	nuclear antigen (EBNA 1)	59	45
427	G00365	Homo sapiens	Human secreted protein, SEQ ID NO: 4446.	99	75
428	AF155819	Mus musculus	doublecortin-like kinase	3448	96
429	Y04315	Homo sapiens	Human secreted protein encoded by gene 23.	385	100
430	AB026891	Homo sapiens	cystine/glutamate transporter	2552	100

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	IDENTITY
431	Y15286	Homo sapiens	vacuolar proton- ATPase subunit M9.2	459	100
432	X81053	Homo sapiens	type IV collagen alpha 4 chain	9706	99
433	U41829	Macaca mulatta	MHC class I antigen Mamu B*07	365	76
434	G03371	Homo sapiens	Human secreted protein, SEQ ID NO: 7452.	100	41
435	AF233238	Gallus gallus	BMP signal transducer Smadl	170	74
436	X52425	Homo sapiens	interleukin 4 receptor	4492	99
437	Y06115	Homo sapiens	Human organic cation transporter OCT-3.	2593	96
438	G02872	Homo sapiens	Human secreted protein, SEQ ID NO: 6953.	130	54
439	L08239	Homo sapiens	located at OATL1	1304	95
440	X17115	Homo sapiens	precursor (AA -15 to 612)	2613	86
441	Y06816	Homo sapiens	Human Notch2 (humN2) protein sequence.	1471	98
442	AB019440	Homo sapiens	immunogloblin heavy chain variable region	545	88
443	¥87350	Homo sapiens	Human signal peptide containing protein HSPP-127 SEQ ID NO:127.	1061	100
444	AJ271736	Homo sapiens	synaptobrevin-like 1 protein	1128	100
445	Y11534	Homo sapiens	PEG1/MEST	1787	100
446	W85719	Homo sapiens	Novel protein (Clone AJ143_1).	271	100
447	Y07900	Homo sapiens	Human secreted protein fragment encoded from gene 49.	87	94
448	X14329	Homo sapiens	carboxypeptidase N precursor (AA -20 to 438)	2463	99
449	M36803	Homo sapiens	hemopexin	2603	100
450	AF116238	Homo sapiens	pseudouridine synthase 1	1927	99
451	AB031051	Homo sapiens	organic anion transporter OATP-E	444	42
452	X16841	Homo sapiens	precursor protein. (-19 to 742)	3958	100
453	AK022830	Homo sapiens	unnamed protein product	373	100

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
454	¥94890	Homo sapiens	Human protein clone HP02798.	637	90
455	AL356014	Arabidopsis thaliana	putative protein	210	38
456	X60221	Homo sapiens	H+-ATP synthase subunit b	1297	99
457	G02532	Homo sapiens	Human secreted protein, SEQ ID NO: 6613.	168	69
458	AJ245375	Homo sapiens	PP35 act	1895	99
459	G00397	Homo sapiens	Human secreted protein, SEQ ID NO: 4478.	57	52
460	AE003708	Drosophila melanogaster	CG6194 gene product	234	65
461	W48352	Homo sapiens	Human breast cancer related protein BCFLT1.	80	60
462	U53420	Rattus norvegicus	sodium-calcium exchanger form 3	397	76
463	Y13402	Homo sapiens	Amino acid sequence of protein PRO310.	1075	63
464	¥27607	Homo sapiens	Human secreted protein encoded by gene No. 41.	610	100
465	L08666	Homo sapiens	porin	122	51
466	¥87084	Homo sapiens	Human secreted protein sequence SEQ ID NO:123.	232	78
467	X16841	Homo sapiens	precursor protein (-19 to 742)	3958	100
468	¥48507	Homo sapiens	Human breast tumour-associated protein 52.	295	91
469	X07973	Ovis aries	MT-Ib protein	84	45
470	W48927	Homo sapiens	Schwannomin-binding protein C-terminal fragment.	78	60
471	AJ224171	Homo sapiens	lipophilin A	454	100
472	G01984	Homo sapiens	Human secreted protein, SEQ ID NO: 6065.	211	64
473	G03793	Homo sapiens	Human secreted protein, SEQ ID NO: 7874.	200	74
474	Y17829	Homo sapiens	Human PRO354 protein sequence.	1006	100
475	Y66706	Homo sapiens	Membrane-bound protein PRO1129.	2153	99
476	G03800	Homo sapiens	Human secreted protein, SEQ ID NO: 7881.	99	78
477	AF216389	Homo sapiens	semaphorin Rs	296	85

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN	IDENTITY
478	X93036	Homo sapiens	MAT8 protein	469	100
479	X53795	Homo sapiens	inducible membrane protein	1412	100
480	AF056195	Homo sapiens	neuroblastoma- amplified protein	4504	98
481	AF116715	Homo sapiens	PRO2829	96	46
482	Z24680	Homo sapiens	garp	167	43
483	Y76198	Homo sapiens	Human secreted protein encoded by gene 75.	82	80
484	AF010144	Homo sapiens	neuronal thread protein AD7c-NTP	324	59
485	Y91592	Homo sapiens	Human secreted protein sequence encoded by gene 6 SEQ ID NO:265.	738	100
486	Y94890	Homo sapiens	Human protein clone HP02798.	605	81
487	U89436	Homo sapiens	tyrosyl-tRNA synthetase	2719	100
488	W88579	Homo sapiens	Secreted protein encoded by gene 46 clone HCFMV39.	479	95
489	G02360	Homo sapiens	Human secreted protein, SEQ ID NO: 6441.	102	70
490	U70976	Homo sapiens	arrestin	1071	61
491	U80746	Homo sapiens	CAGH4	277	81
492	U26361	Helicobacter pylori	Hpn	80	83
493	¥19730	Homo sapiens	SEQ ID NO 448 from WO9922243.	135	53
494	¥27868	Homo sapiens	Human secreted protein encoded by gene No. 107.	185	50
495	AF090901	Homo sapiens	PRO0195	90	46
496	AF061529	Mus musculus	rjs	270	76
497	L34049	Rattus norvegicus	megalin	322	41
498	J04204	Bos taurus	32 kd accessory protein	1743	100
499	Y71118	Homo sapiens	Human Hydrolase protein-16 (HYDRL- 16).	2205	97
500	X13916	Homo sapiens	LDL-receptor related precursor (AA -19 to 4525)	715	92
501	Y00877	Homo sapiens	Human LAPH-2 protein sequence.	138	40
502	¥99368	Homo sapiens	Human PRO1326 (UNQ686) amino acid sequence SEQ ID NO:100.	156	48

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	ક
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE		1		SCORE	
503	Y48308	Homo sapiens	Human prostate	901	100
	İ	1	cancer-associated		
			protein 5.		
504	U67060	Cricetulus	SREBP cleavage	6196	92
		griseus	activating protein		
505	W75857	Homo sapiens	Human secretory	1761	99
		•	protein of clone		
			CO1020-1.		
506	X55764	Homo sapiens	11beta-hydrolase	2604	99
		_	precursor	}	
507	Y41685	Homo sapiens	Human PRO213	1344	94
		_	protein sequence.		
508	X95240	Homo sapiens	cysteine-rich	1368	100
		1	secretory protein-3		
509	AF065482	Homo sapiens	sorting nexin 2	517	77
510	AF135025	Homo sapiens	kallikrein-like	1301	100
			protein 5-related		
			protein 1		
511	AF220492	Homo sapiens	krueppel-like zinc	4100	99
			finger protein HZF2	1200	
512	X58397	Homo sapiens	variable region	670	100
			V251 from V(H)5		200
			qene		
513	W95348	Homo sapiens	Human foetal kidney	406	90
			secreted protein		
			em397_2.		
514	AJ000479	Homo sapiens	putative G-Protein	1966	100
			coupled receptor,	1 - 5 0 0	-00
			EDG6		
515	L05514	Homo sapiens	histatin 3	280	100
516	X95240	Homo sapiens	cysteine-rich	1368	100
			secretory protein-3	-555	
517	D00654	Homo sapiens	enteric smooth	1972	100
			muscle gamma-actin		
518	AJ005453	Mytilus	metallothionein 10	94	35
		edulis	II		1
519	W37864	Homo sapiens	Human protein	362	98
		1	comprising		
			secretory signal		
			amino acid sequence		
			1.		
520	X76091	Homo sapiens	DNA binding protein	3743	99
			RFX2		
521	G03800	Homo sapiens	Human secreted	113	39
· -		- Suproite	protein, SEQ ID NO:		
			7881.		
522	AJ289243	Mus musculus	calpain 12	147	53
523	D30037	Homo sapiens	phosphatidylinosito	1464	100
-	=====		1 transfer protein	7.40.4	1 -00
524	AJ012370	Homo sapiens	NAALADase II	3872	99
•		LIOMO DADIENS	protein	3072	1 22
525	G03909	Homo sapiens	Human secreted	80	41
		LIOMO Saptems	protein, SEQ ID NO:	30	**
	1		Procern' SEG ID MO:		L

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			7990.		
526	U67060	Cricetulus griseus	SREBP cleavage activating protein	6196	92
527	W48351	Homo sapiens	Human breast cancer related protein BCRB2.	85	61
528	AF093408	Homo sapiens	protein kinase A binding protein AKAP110	461	78
529	Y92182	Homo sapiens	Human partial TANGO 195 from clone T195Athpb93f1.	1682	100
530	M28200	Homo sapiens	MHC class II lymphocyte antigen beta chain	432	72
531	X58397	Homo sapiens	variable region V251 from V(H)5 gene	491	74
532	D88577	Mus musculus	Kupffer cell receptor	904	46
533	M84379	Homo sapiens	lymphocyte antigen	1922	97
534	AF279265	Homo sapiens	putative anion transporter 1	212	91
535	AF132035	Homo sapiens	core 2 beta-1,6-N- acetylglucosaminylt ransferase 3	852	92
536	G02958	Homo sapiens	Human secreted protein, SEQ ID NO: 7039.	512	98
537	Y07938	Homo sapiens	Human secreted protein fragment encoded from gene 87.	302	100
538	Y36203	Homo sapiens	Human secreted protein #75.	175	51
539	U16738	Homo sapiens	CAG-isl 7	472	75
540	AL161531	Arabidopsis thaliana	putative proline- rich protein	118	57
541	K00558	Homo sapiens	alpha-tubulin	2393	100
542	U20286	Rattus norvegicus	lamina associated polypeptide 1C	641	55
543	¥27907	Homo sapiens	Human secreted protein encoded by gene No. 119.	128	61
544	AF109674	Rattus norvegicus	late gestation lung protein 1	954	87
545	L35278	Homo sapiens	bone morphogenetic protein	92	40
546	G00541	Homo sapiens	Human secreted protein, SEQ ID NO: 4622.	94	68
			1	+ <u></u>	+
547	AF190664	Mus musculus Homo sapiens	LMBR2 Human 5' EST	113	78

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
NOCEECTIES			secreted protein SEO ID NO:383.	SCORE	
549	AF133816	Homo sapiens	insulin-like peptide INSL5	714	100
550	X70910	Homo sapiens	tetranectin	1069	100
551	M11902	Mus musculus	proline-rich	135	39
			salivary protein		
552	G03477	Homo sapiens	Human secreted protein, SEQ ID NO: 7558.	89	58
553	U63542	Homo sapiens	FAP protein	156	77
554	Y60497	Homo sapiens	Human normal bladder tissue EST encoded protein 169.	89	50
555	Y87303	Homo sapiens	Human signal peptide containing protein HSPP-80 SEQ ID NO:80.	275	100
556	Y17526	Homo sapiens	Human secreted protein clone AM349 2 protein.	1220	100
557	G04064	Homo sapiens	Human secreted protein, SEQ ID NO: 8145.	83	35
558	U51919	Rattus norvegicus	preprocortistatin	84	36
559	AF090901	Homo sapiens	PRO0195	92	66
560	J04031	Homo sapiens	MDMCSF (EC 1.5.1.5; EC 3.5.4.9; EC 6.3.4.3)	226	52
561	AL117237	Homo sapiens	hypothetical protein	4088	94
562	Y50931	Homo sapiens	Human fetal brain cDNA clone vc25_1 derived protein.	485	100
563	Y21631	Homo sapiens	Ligand binding domain of nuclear receptor hTRbeta.	1738	99
564	X90857	Homo sapiens	-14	177	69
565	W35904	Homo sapiens	Human haematopoietic- specific protein (HSP).	862	87
566	W99070	Homo sapiens	Human PIGR-1.	244	90
567	X61653	Homo sapiens	TCR V-beta 13.5	600	100
568	AF166350	Homo sapiens	ST7 protein	4711	99
569	Y07938	Homo sapiens	Human secreted protein fragment encoded from gene 87.	302	100
570	X85019	Homo sapiens	UDP- GalNAc:polypeptide	3069	100

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
			N- acetylgalactosaminy l transferase		
571	U89942	Homo sapiens	lysyl oxidase- related protein	2427	89
572	X04391	Homo sapiens	put. precursor polypeptide	2671	99
573	W36903	Homo sapiens	Human epididymis- specific receptor protein.	5352	100
574	U22816	Homo sapiens	LAR-interacting protein 1b	2042	57
575	Y58618	Homo sapiens	Protein regulating gene expression PRGE-11.	729	57
576	AJ278348	Homo sapiens	pregnancy- associated plasma protein-E	743	100
577	AK024512	Homo sapiens	unnamed protein product	471	100
578	AL031685	Homo sapiens	dJ963K23.4 (KIAA0939 (novel Sodium/hydrogen exchanger family member))	2010	100
579	AF183183	Mus musculus	cochlear otoferlin	116	91
580	W74722	Homo sapiens	Human secreted protein er80_1.	2422	100
581	G03356	Homo sapiens	Human secreted protein, SEQ ID NO: 7437.	. 114	44
582	Y82777	Homo sapiens	Human chordin related protein (Clone dw665_4).	610	98
583	J04988	Homo sapiens	90 kD heat shock protein	3702	100
584	K02576	Homo sapiens	salivary proline- rich protein 1	97	34
585	G03786	Homo sapiens	Human secreted protein, SEQ ID NO: 7867.	159	72
586	AK024490	Homo sapiens	FLJ00092 protein	204	57
587	U22231	Felis catus	ribosomal protein S3a	327	57
588	X55681	Lycopersicon esculentum	extensin (class I)	96	38
589	U68137	Rana ridibunda	prepro-somatostatin	81	33
590	Y19655	Homo sapiens	SEQ ID NO 373 from W09922243.	814	84
591	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	222	56

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
592	AF067801	Homo sapiens	HDCGC21P	116	38
593	X67339	Neurospora	ccg-2	82	37
		crassa	-		
594	G03280	Homo sapiens	Human secreted	169	100
		_	protein, SEQ ID NO:		
			7361.		
595	Y02693	Homo sapiens	Human secreted	130	70
			protein encoded by		
			gene 44 clone		
			HTDAD22.		
596	AE003683	Drosophila	CG9492 gene product	247	56
		melanogaster			
597	Z22968	Homo sapiens	M130 antigen	6205	100
598	AK021847	Homo sapiens	unnamed protein	178	94
599	AP000060		product		<u> </u>
599	AP000060	Aeropyrum pernix	134aa long	80	39
		bernix	hypothetical protein		
600	AK001363	Homo sapiens	unnamed protein	558	92
	ARCOTSOS	nomo saprens	product	336	92
601	G02872	Homo sapiens	Human secreted	147	49
			protein, SEQ ID NO:		"
			6953.		
602	G02538	Homo sapiens	Human secreted	149	65
			protein, SEQ ID NO:		
			6619.		
603	X98330	Homo sapiens	ryanodine receptor	25918	99
			2		
604	AJ243460	Leishmania	proteophosphoglycan	172	35
605	3701000	major			
605	Y81807	Homo sapiens	Human mahogany protein sequence	2499	63
			#2.		
606	AF041069	Equus	fibronectin	109	56
	ATOTIOOS	caballus	libioneccin	109	36
607	Y54591	Homo sapiens	Amino acid sequence	153	77
		1	of a human		
			transferase		
			designated HUTRAN-		
			1.		
608	G03172	Homo sapiens	Human secreted	82	66
			protein, SEQ ID NO:		
			7253.		
609	Y31730	Homo sapiens	Human fused protein	561	99
			kinase-deletion		
			mutant fused C-		
610	Y30163	Homo sapiens	term. Human dorsal root	1112	
010	130103	HOMO Sapiens	receptor 5 hDRR5.	112	49
611	G03714	Homo sapiens	Human secreted	171	70
	303,14	owo pabiens	protein, SEQ ID NO:	1 / 1	/ 0
			7795.		
612	U58514	Homo sapiens	chitinase precursor	402	75
<u> </u>	<u> </u>			<u> </u>	

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
613	AL122105	Homo sapiens	hypothetical protein	399	73
614	AF059198	Homo sapiens	protein kinase/endoribonulc ease	5093	99
615	X17531	Strongylocent rotus purpuratus	epidermal growth factor	234	54
616	AF112982	Homo sapiens	group IID secretory phospholipase A2	852	100
617	AJ006119	Homo sapiens	anti-IFN-G scFv	675	97
618	W54097	Homo sapiens	Homo sapiens B223 sequence.	339	98
619	AF090930	Homo sapiens	PRO0478	141	79
620	W61624	Homo sapiens	Clone HHFEK40 of TM4SF superfamily.	564	98
621	AF119851	Homo sapiens	PRO1722	115	52
622	G03172	Homo sapiens	Human secreted protein, SEQ ID NO: 7253.	173	48
623	¥41379	Homo sapiens	Human secreted protein encoded by gene 72 clone HE6GA29.	261	100
624	U86339	Drosophila grimshawi	expanded	142	36
625	D86853	Catharanthus roseus	extensin	142	39
626	\$58722	Homo sapiens	X-linked retinopathy protein {C-terminal, clone XEH.8c}	116	49
627	G02532	Homo sapiens	Human secreted protein, SEQ ID NO: 6613.	108	50
628	G03790	Homo sapiens	Human secreted protein, SEQ ID NO: 7871.	129	61
629	Y27665	Homo sapiens	Human secreted protein encoded by gene No. 99.	345	100
630	G02837	Homo sapiens	Human secreted protein, SEQ ID NO: 6918.	78	75
631	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	172	65
632	X14329	Homo sapiens	carboxypeptidase N precursor (AA -20 to 438)	2463	99
633	Y87235	Homo sapiens	Human signal peptide containing protein HSPP-12 SEQ	867	100

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN	IDENTITY
NOCHEOTIDE			ID NO:12.	SCORE	ļ
634	W88627	Homo sapiens	Secreted protein	106	73
	1.00027	nome suprem	encoded by gene 94	1 100	/3
			clone HPMBQ32.		
635	W74845	Homo sapiens	Human secreted	395	71
		-	protein encoded by		-
			gene 117 clone		
			HBMUW78.		
636	M16941	Homo sapiens	DR7 beta-chain	1412	100
			glycoprotein		
637	W95634	Homo sapiens	Homo sapiens	1391	100
			secreted protein.		
638	Y78801	Homo sapiens	Hydrophobic domain	1277	100
			containing protein clone HP00631 amino		
			acid sequence.		
639	G03789	Homo sapiens	Human secreted	191	76
		Indiae Edpicins	protein, SEQ ID NO:		/ 0
			7870.		
640	W64535	Homo sapiens	Human leukocyte	2014	99
			cell clone HP00804		1
			protein.		
641	Y94621	Homo sapiens	Epidermal growth	529	91
			factor-like variant		
			in skin-2 amino		
642	G03646	Tions conions	acid sequence. Human secreted	107	
042	G03646	Homo sapiens	protein, SEQ ID NO:	81	42
			7727.		
643	Y87328	Homo sapiens	Human signal	681	100
		•	peptide containing		
			protein HSPP-105		
			SEQ ID NO:105.		
644	Y21386	Homo sapiens	Human HUPF-I mutant	78	31
			protein fragment		
	00000	1	34.		
645	G03790	Homo sapiens	Human secreted protein, SEQ ID NO:	140	55
			7871.		
646	Y35894	Homo sapiens	Extended human	349	100
		Tomo Babiens	secreted protein	347	100
			sequence, SEQ ID		
			NO. 143.		
647	G00517	Homo sapiens	Human secreted	109	37
			protein, SEQ ID NO:		
			4598.		
648	Y25716	Homo sapiens	Human secreted	339	39
			protein encoded		
	5000		from gene 6.		
649	G01246	Homo sapiens	Human secreted	152	80
			protein, SEQ ID NO:		
650	R95913	Homo sapiens	5327. Neural thread	1222	F0
	1193913	nomo sapiens	Mediai chread	233	50

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	0%
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			protein.		
651	Y91469	Homo sapiens	Human secreted protein sequence encoded by gene 19 SEQ ID NO:142.	98	48
652	G03136	Homo sapiens	Human secreted protein, SEQ ID NO: 7217.	94	43
653	U14635	Caenorhabditi s elegans	weak similarity to NADH dehydrogenase	186	30
654	Y14482	Homo sapiens	Fragment of human secreted protein encoded by gene 17.	163	54
655	U14635	Caenorhabditi s elegans	weak similarity to NADH dehydrogenase	186	30
656	AB024565	Mus musculus	heparan sulfate 6- sulfotransferase 2	1128	79
657	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	243	70
658	Y14471	Homo sapiens	Fragment of human secreted protein encoded by gene 4.	95	65
659	AF135381	Homo sapiens	chemokine-like factor 3	89	59
660	U40407	synthetic construct	T cell receptor alpha chain	586	100
661	AF039712	Caenorhabditi s elegans	contains similarity to CDP-alcohol phosphotransferases	289	43
662	G03790	Homo sapiens	Human secreted protein, SEQ ID NO: 7871.	113	55
663	AF084467	Homo sapiens	heparanase	170	32
664	AF279890	Homo sapiens	2P domain potassium channel TREK2	1189	94
665	W63693	Homo sapiens	Human secreted protein 13.	243	84
666	AE003908	Xylella fastidiosa	hypothetical protein	120	28
667	B08948	Homo sapiens	Human secreted protein sequence encoded by gene 21 SEQ ID NO:105.	985	89
668	AF023158	Homo sapiens	tyrosine phosphatase	346	64
669	AF169257	Homo sapiens	sodium/calcium exchanger NCKX3	189	57
670	AF132969	Homo sapiens	CGI-35 protein	364	69
671	AF269286	Homo sapiens	HC6	112	50
672	X98494	Homo sapiens	M phase phosphoprotein 10	529	68
673	G03787	Homo sapiens	Human secreted	83	44
673	G03 /8 /	Homo sapiens	Human secreted	83	44

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	\$ IDENTITY
NOCHEO : IDB			protein, SEQ ID NO: 7868.	SCORE	
674	AF119855	Homo sapiens	PRO1847	123	46
675	AJ242540	Volvox carteri f. nagariensis	hydroxyproline-rich glycoprotein DZ- HRGP	242	42
676	Y91666	Homo sapiens	Human secreted protein sequence encoded by gene 72 SEQ ID NO:339.	529	96
677	Y57936	Homo sapiens	Human transmembrane protein HTMPN-60.	669	100
678	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	156	72
679	W18878	Homo sapiens	Human protein kinase C inhibitor, IPKC-1.	98	68
680	Z12168	Canis familiaris	stimulatory GTP binding protein	980	88
681	G00517	Homo sapiens	Human secreted protein, SEQ ID NO: 4598.	160	48
682	W19932	Homo sapiens	Alzheimer's disease protein encoded by DNA from plasmid pGCS55.	362	100
683	Y30709	Homo sapiens	Amino acid sequence of a human secreted protein.	99	56
684	AF269286	Homo sapiens	HC6	137	72
685	M14362	Homo sapiens	T-cell surface antigen CD2 precursor	275	64
686	G02493	Homo sapiens	Human secreted protein, SEQ ID NO: 6574.	173	61
687	AF248635	Mus musculus	lymphocyte antigen 108 isoform l	303	50
688	D86983	Homo sapiens	similar to D.melanogaster peroxidasin(U11052)	288	55
689	Y59711	Homo sapiens	Secreted protein 58-20-4-G7-FL1.	895	91
690	W48848	Homo sapiens	Human receptor tyrosine kinase LMR3_h N-terminal polypeptide.	1056	89
691	W22652	Homo sapiens	64-863 antibody HSV863 light chain variable region.	459	77
692	AF098066	Homo sapiens	squamous cell carcinoma antigen	1001	98

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	%
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			recognized by T		
693	D83039	Homo sapiens	eti-1	426	98
694	¥79511	Homo sapiens	Human carbohydrate- associated protein CRBAP-7.	1245	99
695	U12623	Rattus norvegicus	cyclic nucleotide gated cation channel	857	83
696	AF229067	Homo sapiens	PADI-H protein	174	61
697	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	196	75
698	U10921	Macaca mulatta	T-cell receptor alpha chain	578	82
699	U31913	Homo sapiens	HBV-X associated protein	167	100
700	X99043	Mus musculus	brain-derived immunoglobulin superfamily molecule	348	82
701	X59770	Homo sapiens	type II interleukin-1 receptor	2130	100
702	AC018758	Homo sapiens	GPI-anchored metastasis- associated protein homolog	207	31
703	Y28816	Homo sapiens	pm4_13 secreted protein.	280	100
704	Y52386	Homo sapiens	Human transmembrane protein HP02000.	1077	100
705	U12392	Haematobia irritans	putative ATPase	481	55
706	U11265	Homo sapiens	HLA-B35	351	92
707	X64594	Homo sapiens	50 kDa erythrocyte plasma membrane glycoprotein	301	88
708	AB046048	Macaca fascicularis	unnamed portein product	260	67
709	G03807	Homo sapiens	Human secreted protein, SEQ ID NO: 7888.	119	60
710	G03315	Homo sapiens	Human secreted protein, SEQ ID NO: 7396.	314	100
711	Y50945	Homo sapiens	Human adult thymus cDNA clone vhl_1 derived protein #1.	742	100
712	G00564	Homo sapiens	Human secreted protein, SEQ ID NO: 4645.	271	98
713	G00125	Homo sapiens	Human secreted	373	80

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	왕
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			protein, SEQ ID NO: 4206.		
714	Y13352	Homo sapiens	Amino acid sequence of protein PRO228.	872	98
715	G02753	Homo sapiens	Human secreted protein, SEQ ID NO: 6834.	222	68
716	Y19588	Homo sapiens	Amino acid sequence of a human secreted protein.	329	100
717	AB030235	Canis familiaris	D4 dopamine receptor	79	35
.718	W74577	Homo sapiens	Human membrane protein BA2303.	748	100
719	Y02693	Homo sapiens	Human secreted protein encoded by gene 44 clone HTDAD22.	235	61
720	X97868	Homo sapiens	arylsulphatase	167	84
721	Y13215	Homo sapiens	Human secreted protein encoded by 5' EST SEQ ID NO: 229.	234	97
722	Y20298	Homo sapiens	Human apolipoprotein E mutant protein fragment 11.	152	39
723	Y86231	Homo sapiens	Human secreted protein HLTHR66, SEQ ID NO:146.	207	51
724	W75083	Homo sapiens	Human secreted protein encoded by gene 27 clone HSPAF93.	685	100
725	W88627	Homo sapiens	Secreted protein encoded by gene 94 clone HPMBQ32.	301	73
726	Y27868	Homo sapiens	Human secreted protein encoded by gene No. 107.	229	58
727	AK025470	Homo sapiens	unnamed protein product	130	64
728	G02872	Homo sapiens	Human secreted protein, SEQ ID NO: 6953.	159	46
729	Y25776	Homo sapiens	Human secreted protein encoded from gene 66.	334	43
730	AF116661	Homo sapiens	PRO1438	153	56
731	W48351	Homo sapiens	Human breast cancer related protein BCRB2.	106	72
732	บ77589	Homo sapiens	MHC class II HLA-	133	69

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	<u> </u>
OF	NUMBER	0120220		WATERMAN	IDENTITY
NUCLEOTIDE	No. Dak			SCORE	
11002201202			DQ-alpha chain		
733	G00357	Homo sapiens	Human secreted	223	67
, 33	300337	nomo bapacino	protein, SEO ID NO:		
			4438.		
734	R28542	Homo sapiens	Human complement	152	96
134	1020342	nome suprem	type 1 receptor		
			SCR9.		
735	Y27868	Homo sapiens	Human secreted	150	65
733	127000	Bapiens	protein encoded by	1 - 3 0	"
			gene No. 107.		
736	AB036706	Homo sapiens	intelectin	368	76
737	Y74042	Homo sapiens	Human prostate	206	65
131	1/4042	nomo sapiens	tumor EST fragment	200	63
			derived protein		
			#229.	ł	
720	152 63 5 6	******	Human secreted	153	77
738	Y36156	Homo sapiens		153	' '
	125 4 2 2 2		protein #28.	1751	79
739	W74802	Homo sapiens	Human secreted	1/51	19
			protein encoded by		
			gene 73 clone		
			HSQEL25.		
740	W85614	Homo sapiens	Secreted protein	224	91
			clone fr473_2.		
741	Y13377	Homo sapiens	Amino acid sequence	394	98
			of protein PRO257.		
742	Z69384	Caenorhabditi	Similarity to	515	45
		s elegans	Salmonella	ł	Ì
			regulatory protein	1	
			UHPC		
			(SW:UHPC_SALTY)		
743	W47589	Homo sapiens	T-cell receptor	681	92
	<u>L</u> .		beta-chain.		
744	G03786	Homo sapiens	Human secreted	243	71
			protein, SEQ ID NO:		
			7867.		
745	Y50690	Homo sapiens	Human Hum4 VL ClaI-	540	81
			HindIII segment		
			encoded protein.	Ĺ	
746	U03414	Rattus	neuronal	363	67
		norvegicus	olfactomedin-		
			related ER		
	1 .		localized protein		1
747	G00352	Homo sapiens	Human secreted	84	51
			protein, SEQ ID NO:		
	1		4433.	1	1
748	Y02671	Homo sapiens	Human secreted	145	60
			protein encoded by		
			gene 22 clone		
	1		HMSJW18.		
749	AF026919	Homo sapiens	amyloid lambda	557	83
	31 020919	Paprens	light chain	1	33
			variable region	1	
750	X76732	Home geniene	NEFA protein	297	100
130	A/0/34	Homo sapiens	MELY Proceru	1 431	1 100

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	용
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
751	R92754	Homo sapiens	Human growth differentiation factor-12.	628	100
752	Y91462	Homo sapiens	Human secreted protein sequence encoded by gene 12	597	100
753	Y66700	Homo sapiens	SEQ ID NO:135. Membrane-bound protein PRO1137.	754	99
754	G01648	Homo sapiens	Human secreted protein, SEQ ID NO: 5729.	281	100
755	AB040434	Homo sapiens	hTROY	752	100
756	Y28680	Homo sapiens	Human nm214_3 secreted protein.	178	44
757	W75100	Homo sapiens	Human secreted protein encoded by gene 44 clone HE8CJ26.	203	66
758	AF090930	Homo sapiens	PRO0478	87	45
759	D84336	Rattus norvegicus	ZOG	484	48
760	W88627	Homo sapiens	Secreted protein encoded by gene 94 clone HPMBQ32.	150	81
761	Y48616	Homo sapiens	Human breast tumour-associated protein 77.	569	70
762	Y87320	Homo sapiens	Human signal peptide containing protein HSPP-97 SEQ ID NO:97.	918	100
763	G03655	Homo sapiens	Human secreted protein, SEQ ID NO: 7736.	248	89
764	AF031174	Homo sapiens	Ig-like membrane protein	428	45
765	U08255	Rattus norvegicus	glutamate receptor delta-1 subunit	802	99
766	¥99369	Homo sapiens	Human PR01249 (UNQ632) amino acid sequence SEQ ID NO:102.	4578	99
767	AK001586	Homo sapiens	unnamed protein product	973	98
768	AC007063	Arabidopsis thaliana	putative ABC transporter	126	31
769	AF303378	Homo sapiens	sialic acid- specific acetylesterase II	713	100
770	G00517	Homo sapiens	Human secreted protein, SEQ ID NO: 4598.	90	37

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
771	Y59733	Homo sapiens	Human normal	1253	99
			ovarian tissue		
	3.77.20056	 	derived protein 10.	1.53	-
772	AF132856	Homo sapiens	suppressor of G2 allele of skp1	163	86
			homolog	i	
773	AB029482	Mus musculus	JNK-binding protein	1082	97
773	AB029402	rius muscurus	JNKBP1		3 /
774	G02108	Homo sapiens	Human secreted	134	62
			protein, SEQ ID NO:		
			6189.		
775	AB047818	Homo sapiens	Soggy	1239	100
776	Y66689	Homo sapiens	Membrane-bound	804	99
	120000		protein PRO1136.		
777	¥71107	Homo sapiens	Human Hydrolase protein-5 (HYDRL-	733	99
			protein-5 (HYDRL-		
778	AC005626	Homo sapiens	R29124 1	182	38
779	W88707	Homo sapiens	Secreted protein	126	56
1,13	W00707	nomo sapiens	encoded by gene 174	123	
			clone HE9FB42.		
780	G03657	Homo sapiens	Human secreted	455	96
		•	protein, SEQ ID NO:		
			7738.		
781	AJ001616	Mus musculus	myeloid associated	201	36
:			differentiation		
			protein		
782	Y64942	Homo sapiens	Human 5' EST	86	65
]			related polypeptide SEO ID NO:1103.		
783	AL356276	Warra anni anna	bA367J7.2.1 (novel	845	91
/63	ALI336276	Homo sapiens	Immunoglobulin	045	1 31
			domains containing		
			protein (isoform		•
			1))		
784	Y00876	Homo sapiens	Human LAPH-1	291	43
			protein sequence.		
785	G00270	Homo sapiens	Human secreted	603	100
			protein, SEQ ID NO:		
			4351.		
786	AF154121	Homo sapiens	sodium-dependent	864	100
			high-affinity		
			dicarboxylate		
707	720004	172	transporter Human GABA B	ļ	42
787	Y29804	Homo sapiens	receptor subunit	83	42
			HG20 peptide #6.		1
788	AL080239	Homo sapiens	bG256022.1 (similar	599	100
, 50	AU000233	nomo saprens	to IGFALS (insulin-	"	100
			like growth factor		
			binding protein,		
			acid labile		
			subunit))		J

TABLE 2

SEQ ID NO: OF	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN	% IDENTITY
NUCLEOTIDE 789	77.021.056			SCORE	
789	AL031856	Schizosacchar omyces pombe	PUTATIVE GOLGI URIDINE DIPHOSPHATE-N- ACETYLGLUCOSAMINE TRANSPORTER	192	40
790	G03448	Homo sapiens	Human secreted protein, SEQ ID NO: 7529.	141	43
791	U81291	Xenopus laevis	oviductin	310	38
792	Y41332	Homo sapiens	Human secreted protein encoded by gene 25 clone HPIBO48.	295	50
793	L20315	Mus musculus	MPS1 protein	702	77
794	G01314	Homo sapiens	Human secreted protein, SEQ ID NO: 5395.	91	36
795	AF003136	Caenorhabditi s elegans	similar to 1-acyl- glycerol-3- phosphate acyltransferases	122	38
796	G00637	Homo sapiens	Human secreted protein, SEQ ID NO: 4718.	160	67
797	Y36144	Homo sapiens	Human secreted protein #16.	622	100
798	U09453	Cricetulus griseus	UDP-N- acetylglucosamine: dolichyl phosphate N-acetylglucosamine 1-phosphate transferase	178	66
799	Y76144	Homo sapiens	Human secreted protein encoded by gene 21.	633	100
800	Y73456	Homo sapiens	Human secreted protein clone yd145_1 protein sequence SEQ ID NO:134.	413	89
801	¥86540	Homo sapiens	Human gene 77- encoded protein fragment, SEQ ID NO:457.	443	96
802	U 49973	Homo sapiens	ORF1; MER37; putative transposase similar to pogo element	311	53
803	M63573	Homo sapiens	secreted cyclophilin-like protein	700	88
804	AF091622	Homo sapiens	PHD finger protein	177	100

TABLE 2

SEQ ID NO: OF	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	
805	W37869	Homo sapiens	Human protein	381	100
			comprising secretory signal amino acid sequence 6.		
806	G03556	Homo sapiens	Human secreted protein, SEQ ID NO: 7637.	221	72
807	AF178941	Homo sapiens	ATP-binding cassette sub-family A member 2	583	87
808	Y91385	Homo sapiens	Human secreted protein sequence encoded by gene 40 SEQ ID NO:106.	786	100
809	Y00826	Rattus norvegicus	gp210 (AA 1-1886)	169	83
810	G03143	Homo sapiens	Human secreted protein, SEQ ID NO: 7224.	328	100
811	W00870	Homo sapiens	Polycystic kidney disease 1 (PKD1) polypeptide.	22446	99
812	¥73434	Homo sapiens	Human secreted protein clone yd51_1 protein sequence SEQ ID NO:90.	417	90
813	AB031996	Ralstonia sp. KN1	ferredoxin-like protein	94	44
814	AF201734	Mus musculus	testis specific serine kinase-3	800	87
815	Y01181	Homo sapiens	Polypeptide fragment encoded by gene 12.	68	55
816	Y76166	Homo sapiens	Human secreted protein encoded by gene 43.	724	94
817	AL109827	Homo sapiens	dJ309K20.2 (acrosomal protein ACR55 (similar to rat sperm antigen 4 (SPAG4)))	639	84
818	M62829	Homo sapiens	ETR103	137	53
819	Y38422	Homo sapiens	Human secreted protein.	526	100
820	AF119815	Homo sapiens	G-protein-coupled receptor	561	79
821	Y87101	Homo sapiens	Human secreted protein sequence SEQ ID NO:140.	628	100
822	M91463	Homo sapiens	glucose transporter	213	79

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
823	L34938	Rattus	ionotropic	618	90
		norvegicus	glutamate receptor		
824	W17846	Homo sapiens	Cytosolic phospholipase A2/B (clone 19b product).	209	64
825	Y66722	Homo sapiens	Membrane-bound protein PRO1104.	221	67
826	G02493	Homo sapiens	Human secreted protein, SEQ ID NO: 6574.	138	72
827	Y91423	Homo sapiens	Human secreted protein sequence encoded by gene 11 SEQ ID NO:144.	671	54
828	U78090	Rattus norvegicus	potassium channel regulator 1	502	80
829	U08813	Oryctolagus cuniculus	597 aa protein related to Na/glucose cotransporters	906	84
830	AJ272063	Homo sapiens	vanilloid receptor	630	90
831	U36898	Rattus norvegicus	pheromone receptor VN6	135	52
832	Z46973	Homo sapiens	phosphatidylinosito 1 3-kinase	396	80
833	Y95433	Homo sapiens	Human calcium channel SOC-2/CRAC-1 C-terminal polypeptide.	747	99
834	AF132856	Homo sapiens	suppressor of G2 allele of skp1 homolog	163	86
835	AC006042	Homo sapiens	supported by human ESTs AI681256.1(NID:g489 1438),N32168.1(NID: g1152567), and genscan	195	87
836	B01247	Homo sapiens	Human HE6 receptor.	371	45
837	G03788	Homo sapiens	Human secreted protein, SEQ ID NO: 7869.	196	59
838	U70136	Homo sapiens	megakaryocyte stimulating factor; MSF	6954	98
839	AF017153	Mus musculus	putative RNA helicase and RNA dependent ATPase	178	51
840	Y31830	Homo sapiens	Human adult brain secreted protein nh899 8.	244	56

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	<u> </u>
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	-
841	Y27593	Homo sapiens	Human secreted	437	81
			protein encoded by		
		1	gene No. 27.	l	1.
842	G01984	Homo sapiens	Human secreted	196	74
			protein, SEQ ID NO:		}
			6065.		
843	AL008723	Homo sapiens	dJ90G24.4 (SAAT1	183	92
			(low affinity		ļ
			sodium glucose cotransporter		
			cotransporter (sodium:solute		
•			symporter family)))]
844	AF068065	Cryptosporidi	GP900: mucin-like	263	47
044	AFOSSOCS	um parvum	glycoprotein	203	* '
845	Y00815	Homo sapiens	put. LAR preprotein	341	100
043	100015	nomo saprens	(AA -16 to 1881)	712	
846	Y06816	Homo sapiens	Human Notch2	1224	99
• • • • • • • • • • • • • • • • • • • •			(humN2) protein		
			sequence.		
847	AF104923	Homo sapiens	putative	293	95
	1		transcription		
			factor		
848	Y09945	Rattus	putative integral	589	53
•		norvegicus	membrane transport		
			protein		
849	AL157874	Schizosacchar	similar to yeast	146	40
		omyces pombe	SCT1 suppressor of a choline transport		
			mutant		
850	R71003	Homo sapiens	Human neuronal	141	89
030	1272005	nome Bupiens	calcium channel	***	
			subunit alpha 1c-1.		
851	X75756	Homo sapiens	protein kinase C mu	318	90
852	AF142676	Drosophila	sodium-hydrogen	366	48
		melanogaster	exchanger NHE1		
853	Y45381	Homo sapiens	Human secreted	139	73
			protein fragment	İ	
			encoded from gene		
			28.		
854	G03789	Homo sapiens	Human secreted	121	60
			protein, SEQ ID NO:		
0.55	7765400	177	7870.	1 100	
855	U65409	Yarrowia lipolytica	Sla2p	109	25
856	M19419	Mus musculus	proline-rich	109	36
0.20	1/11/24/12	rus muscurus	salivary protein	109	30
857	Y99355	Homo sapiens	Human PRO1295	667	98
<i>55</i> ,	1,5555	Dapiens	(UNQ664) amino acid		-
			sequence SEQ ID		
			NO:54.		
858	W19919	Homo sapiens	Human Ksr-1 (kinase	211	86
			suppressor of Ras).		

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION .	SMITH- WATERMAN SCORE	% IDENTITY
			channel SOC-3/CRAC-2.		
860	AF070066	Mus musculus	Citron-K kinase	628	97
861	AF286095	Homo sapiens	IL-22 receptor	933	100
862	AF020195	Mus musculus	pancreas sodium bicarbonate cotransporter	475	68
863	G03712	Homo sapiens	Human secreted protein, SEQ ID NO: 7793.	240	100
864	AF195092	Homo sapiens	sialic acid-binding immunoglobulin-like lectin-8	288	87
865	AF208110	Homo sapiens	IL-17 receptor homolog precursor	2688	99
866	L42338	Mus musculus	sodium channel 25	733	98
867	G02360	Homo sapiens	Human secreted protein, SEQ ID NO: 6441.	101	70
868	AF065215	Homo sapiens	cytosolic phospholipase A2 beta	290	42
869	L43631	Homo sapiens	scaffold attachment factor B	106	95
870	G03034	Homo sapiens	Human secreted protein, SEQ ID NO: 7115.	108	54
871	Z21514	Rattus norvegicus	integral membrane glycoprotein	84	47
872	AF097518	Homo sapiens	liver-specific transporter	147	40
873	AF288223	Drosophila melanogaster	Crossveinless 2	136	39
874	U90126	Bos taurus	ABC transporter	245	36
875	AF099988	Mus musculus	Ste-20 related kinase SPAK	103	34
876	¥70400	Homo sapiens	Human cell- signalling protein- 2.	220	86
877	¥36300	Homo sapiens	Human secreted protein encoded by gene 77.	1863	99
878	AF151074	Homo sapiens	HSPC240	193	29
879	Y94951	Homo sapiens	Human secreted protein clone dw78_1 protein sequence SEQ ID NO:108.	251	89
880	AF165310	Homo sapiens	ATP cassette binding transporter 1	231	31
881	AF252281	Mus musculus	Kelch-like 1 protein	256	58

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF	NUMBER	}		WATERMAN	IDENTITY
NUCLEOTIDE				SCORE	<u> </u>
882	Y00931	Homo sapiens	Prostate-tumour	1039	98
			derived antigen #4.		
883	Y27576	Homo sapiens	Human secreted	394	96
			protein encoded by		
004	U00009	Escherichia	gene No. 10.	153	30
884	000009	coli	yeer	123	30
885	Y57945	Homo sapiens	Human transmembrane protein HTMPN-69.	1543	100
886	Y28678	Homo sapiens	Human cw272_7 secreted protein.	375	60
887	W95349	Homo sapiens	Human foetal brain secreted protein fh170 7.	377	89
888	Y87329	Homo sapiens	Human signal peptide containing protein HSPP-106 SEO ID NO:106.	285	89
889	AL121845	Homo sapiens	dJ583P15.5.1 (novel protein (isoform 1))	1399	99
890	R75181	Homo sapiens	Partial peptide of human HMW kininogen fragment 1.2.	100	29
891	AF105365	Homo sapiens	K-Cl cotransporter KCC4	680	100
892	Y91644	Homo sapiens	Human secreted protein sequence encoded by gene 43 SEQ ID NO:317.	673	95
893	S52051	Rattus sp.	neurotransmitter transporter	656	99
894	\$52051	Rattus sp.	neurotransmitter transporter	617	94
895	R47120	Homo sapiens	Partial human H13 polypeptide.	343	60
896	Z98046	Homo sapiens	dJ1409.2 (Melanoma- Associated Antigen MAGE LIKE)	332	49
897	AJ006203	Oryctolagus cuniculus	capacitative calcium entry channel 2	740	99
898	AF156547	Mus musculus	putative E1-E2 ATPase	769	95
899	AC004076	Homo sapiens	R30217_1	788	98
900	D00099	Homo sapiens	Na,K-ATPase alpha- subunit	753	94
901	R27648	Homo sapiens	Human calcium channel 27980/10.	536	85
902	Y57955	Homo sapiens	Human transmembrane protein HTMPN-79.	606	100
903	AF155913	Mus musculus	putative E1-E2 ATPase	1039	85

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	\$ IDENTITY
904	Y73446	Homo sapiens	Human secreted protein clone yc27_1 protein sequence SEQ ID NO:114.	369	66
905	Y94903	Homo sapiens	Human secreted protein clone pt332_1 protein sequence SEQ ID NO:12.	3777	100
906	AB032470	Homo sapiens	seven transmembrane protein TM7SF3	2124	100
907	G00517	Homo sapiens	Human secreted protein, SEQ ID NO: 4598.	90	50
908	AF010144	Homo sapiens	neuronal thread protein AD7c-NTP	270	65
909	AF263912	Streptomyces noursei	NysA	113	25
910	Y53051	Homo sapiens	Human secreted protein clone dd119_4 protein sequence SEQ ID NO:108.	843	49
911	Y76179	Homo sapiens	Human secreted protein encoded by gene 56.	634	100
912	G00352	Homo sapiens	Human secreted protein, SEQ ID NO: 4433.	229	71
913	U93569	Homo sapiens	p40	110	32
914	G02639	Homo sapiens	Human secreted protein, SEQ ID NO: 6720.	65	46
915	Y94951	Homo sapiens	Human secreted protein clone dw78_1 protein sequence SEQ ID NO:108.	100	38
916	G03263	Homo sapiens	Human secreted protein, SEQ ID NO: 7344.	80	47
917	W74887	Homo sapiens	Human secreted protein encoded by gene 160 clone HCELB21.	273	69
918	¥73464	Homo sapiens	Human secreted protein clone yl4_1 protein sequence SEQ ID NO:150.	982	90
919	AF064801	Homo sapiens	multiple membrane spanning receptor TRC8	551	32

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	\$ IDENTITY
920	¥87335	Homo sapiens	Human signal peptide containing protein HSPP-112 SEQ ID NO:112.	622	99
921	AK000496	Homo sapiens	unnamed protein product	342	74
922	Y41360	Homo sapiens	Human secreted protein encoded by gene 53 clone HJPAD75.	367	100
923	G02872	Homo sapiens	Human secreted protein, SEQ ID NO: 6953.	328	75
924	Y53881	Homo sapiens	A suppressor of cytokine signalling protein designated HSCOP-1.	1489	100
925	AC004144	Homo sapiens	R34001_1	193	60
926	AF119851	Homo sapiens	PRO1722	153	82
927	G02654	Homo sapiens	Human secreted protein, SEQ ID NO: 6735.	82	57
928	Y30819	Homo sapiens	Human secreted protein encoded from gene 9.	264	33
929	G01691	Homo sapiens	Human secreted protein, SEQ ID NO: 5772.	66	43
930	AF187845	Homo sapiens	small protein effector 1 of Cdc42	431	100
931	AL390114	Leishmania major	extremely cysteine/valine rich protein	113	40
932	AL080239	Homo sapiens	bG256022.1 (similar to IGFALS (insulin-like growth factor binding protein, acid labile subunit))	1451	97
933	W85613	Homo sapiens	Secreted protein clone fm60_1.	234	100
934	AF009243	Homo sapiens	proline-rich Gla protein 2	223	42
935	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	271	66
936	AK000385	Homo sapiens	unnamed protein product	193	64
937	AF010144	Homo sapiens	neuronal thread protein AD7c-NTP	270	65
938	AF119851	Homo sapiens	PRO1722	170	71
939	Y07922	Homo sapiens	Human secreted protein fragment	226	95

TABLE 2

SEQ ID NO: OF NUCLEOTIDE	ACCESSION NUMBER	SPECIES	DESCRIPTION	SMITH- WATERMAN SCORE	% IDENTITY
NOCHECTIES	 		encoded from gene	SCORE	
940	Y41712	Homo sapiens	Human PRO724 protein sequence.	653	96
941	AF010144	Homo sapiens	neuronal thread protein AD7c-NTP	310	64
942	Y45318	Homo sapiens	Human secreted protein fragment encoded from gene	502	98
943	Y07899	Homo sapiens	Human secreted protein fragment encoded from gene 48.	309	98
944	X92485	Plasmodium vivax	pval	185	51
945	AJ289133	Mus musculus	chondroitin 4-0- sulfotransferase	565	43
946	AF151074	Homo sapiens	HSPC240	1337	99
947	U40829	Saccharomyces cerevisiae	Weak similarity near C-terminus to RNA Polymerase beta subunit (Swiss Prot. accession number P11213) and CCAAT-binding transcription factor (PIR accession number A36368)	361	50
948	¥87285	Homo sapiens	Human signal peptide containing protein HSPP-62 SEQ ID NO:62.	348	82
949	Y86230	Homo sapiens	Human secreted protein HKFBC53, SEQ ID NO:145.	368	80
950	AJ010346	Homo sapiens	RING-H2	333	87
951	Z56281	Homo sapiens	interferon regulatory factor 3	1573	81
952	Y57896	Homo sapiens	Human transmembrane protein HTMPN-20.	421	100
953	G03789	Homo sapiens	Human secreted protein, SEQ ID NO: 7870.	135	55
954	Y87103	Homo sapiens	Human secreted protein sequence SEQ ID NO:142.	83	50
955	Y87345	Homo sapiens	Human signal peptide containing protein HSPP-122 SEQ ID NO:122.	885	99
956	X81479	Homo sapiens	EMR1	1148	99

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	olo
OF	NUMBER			WATERMAN SCORE	IDENTITY
NUCLEOTIDE	377777			4061	99
957	AF175406	Homo sapiens	transient receptor potential 4	4061	99
958	G03789	Homo sapiens	Human secreted	276	73
958	G03/89	Homo sapiens	protein, SEQ ID NO:	276	′³
			7870.		
959	M63274	Plasmodium	malaria antigen	77	38
959	M632/4	falciparum	maiaria ancigen	' '	30
960	¥78795	Homo sapiens	Human antizuai-2	3384	83
960	1/6/95	MOMO Sapiens	(AZ-2) amino acid	3304	03
			sequence.		1
961	AL133469	Streptomyces	putative secreted	139	41
961	ALLISTED	coelicolor	proline-rich		
		A3 (2)	protein		
962	G03787	Homo sapiens	Human secreted	232	72
902	903787	nomo saprens	protein, SEQ ID NO:	1 2 2	
			7868.		
963	W74828	Homo sapiens	Human secreted	1016	99
903	W/4020	nomo saprens	protein encoded by	1 -020	
			gene 100 clone		
			HLOAB52.		
964	W48351	Homo sapiens	Human breast cancer	226	58
204	"40331	nomo supromo	related protein		
			BCRB2.		
965	X63893	Sus scrofa	alpha-stimulatory	319	86
505	1.00003		subunit of GTP-		
			binding protein		
966	AB033019	Homo sapiens	KIAA1193 protein	245	97
967	Y36156	Homo sapiens	Human secreted	223	85
		_	protein #28.		
968	AF119851	Homo sapiens	PRO1722	188	69
969	Y15224	Homo sapiens	Human receptor	214	42
		_	protein (HURP) 3		İ
			amino acid		
			sequence.		
970	G02754	Homo sapiens	Human secreted	81	62
			protein, SEQ ID NO:		
			6835.		
971	U22376	Homo sapiens	alternatively	212	81
	,	_	spliced product		
			using exon 13A		
972	W74870	Homo sapiens	Human secreted	164	81
		- '	protein encoded by		
			gene 142 clone		
	1		HTWCB92.		
973	Y30817	Homo sapiens	Human secreted	717	98
			protein encoded		1
			from gene 7.		
974	AF079529	Homo sapiens	cAMP-specific	2353	96
			phosphodiesterase		
			8B; PDE8B1; 3',5'-		1
			cyclic nucleotide	1	
r.			phosphodiesterase		
975	AF099028	Drosophila	putative	1061	52
	1-11-055020		F		<u> </u>

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	%
OF	NUMBER			WATERMAN	IDENTITY
NUCLEOTIDE	<u> </u>			SCORE '	
		melanogaster	transmembrane		
0.00			protein cmp44E	<u> </u>	
976	G03786	Homo sapiens	Human secreted	179	72
			protein, SEQ ID NO: 7867.		
977	Y22495	Homo sapiens	Human secreted	1629	100
911	122495	nomo saprens	protein sequence	1629	100
			clone ch4 11.		
978	W74813	Homo sapiens	Human secreted	722	92
2,70		Tromo bapacino	protein encoded by	1 122	122
			gene 85 clone		
			HSDFV29.		
979	AK023408	Homo sapiens	unnamed protein	974	96
			product	1 - / -	
980	AF229178	Homo sapiens	leucine rich repeat	276	67
			and death domain		
			containing protein		
981	G03797	Homo sapiens	Human secreted	198	56
		1	protein, SEQ ID NO:		
			7878.		
982	W74831	Homo sapiens	Human secreted	153	100
			protein encoded by		
			gene 103 clone		
			HEBDJ82.		[
983	G01335	Homo sapiens	Human secreted	157	96
			protein, SEQ ID NO:		
			5416.		
984	Y73436	Homo sapiens	Human secreted	450	100
			protein clone		
			ye43_1 protein		
			sequence SEQ ID NO:94.		
985	G00354	Tieme genieng	Human secreted	96	
905	G00354	Homo sapiens	protein, SEQ ID NO:	96	58
			4435.		
986	Y41712	Homo sapiens	Human PRO724	639	88
	111/11	nomo saprens	protein sequence.	033	
987	Y57896	Homo sapiens	Human transmembrane	421	100
			protein HTMPN-20.	12	100
988	Y66691	Homo sapiens	Membrane-bound	716	65
			protein PRO809.	1	
989	AF090943	Homo sapiens	PRO0659	926	100
990	G00403	Homo sapiens	Human secreted	80	46
		•	protein, SEQ ID NO:		
			4484.		
991	G03411	Homo sapiens	Human secreted	62	57
		_	protein, SEQ ID NO:		
		1	7492.		
992	G00270	Homo sapiens	Human secreted	143	96
			protein, SEQ ID NO:		
		1	4351.		
993	AF026246	Homo sapiens	HERV-E integrase	361	80
994	Y36421	Homo sapiens	Fragment of human	83	37

TABLE 2

SEQ ID NO:	ACCESSION	SPECIES	DESCRIPTION	SMITH-	8
OF NUCLEOTIDE	NUMBER			WATERMAN SCORE	IDENTITY
			secreted protein encoded by gene 8.		
995	U22376	Homo sapiens	alternatively spliced product using exon 13A	175	78
996	G03790	Homo sapiens	Human secreted protein, SEQ ID NO: 7871.	87	35
997	G00397	Homo sapiens	Human secreted protein, SEQ ID NO: 4478.	149	61
998	J02642	Homo sapiens	glyceraldehyde 3- phosphate dehydrogenase (EC 1.2.1.12)	429	69
999	AF119851	Homo sapiens	PRO1722	204	50
1000	¥91423	Homo sapiens	Human secreted protein sequence encoded by gene 11 SEQ ID NO:144.	393	53
1001	Y66695	Homo sapiens	Membrane-bound protein PRO1344.	1183	87
1002	AF090931	Homo sapiens	PRO0483	149	68
1003	Y33261	Homo sapiens	Human p99 protein.	314	59
1004	U11494	Mus musculus	protein kinase	360	77
1005	AK021848	Homo sapiens	unnamed protein product	186	69
1006	Y13892	Homo sapiens	PI-3 kinase	233	97
1007	W48351	Homo sapiens	Human breast cancer related protein BCRB2.	144	65
1008	G03793	Homo sapiens	Human secreted protein, SEQ ID NO: 7874.	202	67
1009	U91682	Aedes aegypti	vitelline membrane protein homolog	88	42

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		,	REGION	REGION
1	1010	100	299	535
2	1011	1002	19	267
3	1012	1003	31	423
4	1013	1007	148	840
5	1014	1009	139	318
6	1015	1010	413	748
7	1016	1012	357	154
8	1017	1014	133	285
9	1018	1016	61	441
10	1019	102	269	832
11	1020	1021	148	342
12	1021	1022	45.	452
13	1022	1035	222	779
14	1023	1038	222	779
15	1024	1042	735	517
16	1025	1049	120	320
17	1026	1055	195	395
18	1027	1061	13	189
19	1028	1070	972	1109
20	1029	1071	1504	1686
21	1030	1077	425	574
22	1031	108	46	501
23	1032	1088	1949	7240
24	1033	1092	119	571
25	1034	1095	118	564
26	1035	1096	110	373
27	1036	1098	66	353
28	1037	1099	1	417
29	1038	11	764	573
30	1039	1100	157	1014
31	1040	1102	1526	1813
32	1041	1103	1529	1338
33	1042	1104	685	1929
34	1043	1105	887	744
35	1044	1110	880	443
36	1045	1111	696	538
37	1046	1113	52	1272
38	1047	1117	1357	554
39	1048	1118	1478	1654
40	1049	112	482	712
41	1050	1121	3	1424
42	1051	1130	131	271
43	1052	1132	849	151
44	1053	1137	265	705
45	1054	1138	13	381
46	1055	1140	51	416
47	1056	1146	2389	2541
48	1057	1148	1517	738
49	1058	115	179	334
50	1059	1154	68	358
	<u> </u>	<u> </u>		I

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
NOCHEOTIDE	ACID	05/451,404	REGION	REGION
51	1060	1155	34	330
52	1061	1157	242	433
53	1062	1160	410	856
54	1063	1161	154	342
55	1064	1163	202	477
56	1065	1167	72	272
57	1066	117	235	2
58	1067	1170	47	211
59	1068	1176	16	159
60	1069	1177	135	326
61	1070	118	1248	1466
62	1071	1183	431	886
63	1072	1187	191	529
	1072		1303	1148
64		1189	380	613
65	1074	119		1272
66	1075	1190	514	
67	1076	1192	1529	1338
68	1077	1197	93	533
69	1078	1199	227	391
70	1079	1202	117	407
71	1080	1204	12	413
72	1081	1205	49	603
73	1082	1216	487	1341
74	1083	1217	982	764
75	1084	1228	99	266
76	1085	1230	973	770
77	1086	1233	233	418
78	1087	1234	2959	2078
79	1088	1235	112	1542
80	1089	1239	3019	2822
81	1090	1242	1335	781
82	1091	1248	29	169
83	1092	125	542	405
84	1093	1250	1381	1572
85	1094	1252	480	226
86	1095	1255	19	285
87	1096	1259	165	638
88	1097	126	627	364
89	1098	1260	289	462
90	1099	1262	138	353
91	1100	1264	1159	1299
92	1101	1266	13	402
93	1102	1269	296	805
94	1103	127	212	397
95	1104	1270	126	374
96	1105	1272	2025	2396
97	1105	1273	1367	624
98	1107	1274	1108	746
99	1107	1274	919	1077
100				1272
100	1109	1279	496	14/4

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		05,151,101	REGION	REGION
101	1110	1283	265	125
102	1111	1287	107	385
103	1112	1297	333	545
104	1113	13	187	47
105	1114	130	126	290
106	1115	1306	323	75
107	1116	1308	457	891
108	1117	1311	258	674
109	1118	1315	242	823
110	1119	1317	82	435
111	1120	1319	781	3306
112	1121	1323	1402	1671
113	1122	1329	279	665
114	1123	1336	37	765
115	1124	1337	177	389
116	1125	1338	887	744
117	1126	1339	248	724
118	1127	1341	298	525
119	1128	1342	26	445
120	1129	1344	23	
121	1130	1345	160	370
122	1131	1351	2737	402
123	1132	1353	655	2600
124	1133	1354	94	792
125	1134	1354	679	354
126	1135	1358	679	849
127	1136	1359	32	849
128	1137	1361	271	346
129	1138	1362	637	426
130	1139	1363		1197
131	1140	1364	24	350
132	1141	1364	119	367
133	1142		111	284
134	1142	1377	1221	1358
135	1144	1378	643	470
136		<u> </u>	99	539
137	1145	1382	994	686
138	1146 1147	1384	34	264
139		1386	124	477
140	1148	1389	1197	1
	1149	139	94	294
141	1150	1390	1262	1053
142	1151	1393	1182	1325
143	1152	1394	1351	1542
144	1153	1395	229	411
145	1154	1396	923	1147
146	1155	1397	49	252
147	1156	1398	684	863
148	1157	1399	2613	286
149	1158 1159	14	997	758
150				

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
ROCEEGIEDE	1.022	05, 151, 101	REGION	REGION
151	1160	1406	735	1235
152	1161	1407	967	716
153	1162	1408	75	314
154	1163	1409	101	313
155	1164	141	384	551
156	1165	1414	242	532
157	1166	142	158	15
158	1167	1421	604	1425
159	1168	1422	1146	1835
160	1169	1423	2657	3295
161	1170	1424	315	163
162	1171	1426	39	509
163	1172	1427	892	686
164	1173	1428	395	619
165	1174	1430	284	514
166	1175	1432	178	2
167	1176	1433	1136	972
168	1177	1435	1283	1540
169	1178	1436	1669	2235
170	1179	144	55	219
171	1180	1440	363	121
172	1181	1441	1991	2197
173	1182	1443	1765	3054
174	1183	1445	1023	865
175	1184	1446	5692	5859
176	1185	1447	2959	2078
177	1186	1448	775	945
178	1187	1451	858	1430
179	1188	1453	1370	723
180	1189	1455	480	1007
181	1190	1457	278	451
182	1191	1459	824	561
183	1192	1460	56	463
184	1193	1461	184	480
185	1194	1462	486	635
186	1195	1465	319	492
187	1196	1466	398	3
188	1197	1468	262	453
189	1198	1476	526	684
190	1199	148	271	420
191	1200	1482	568	714
192	1201	1484	203	340
193	1202	1486	2185	1190
194	1203	1492	438	2912
195	1204	1493	82	225
196	1205	1501	210	347
197	1206	1508	1364	1101
198	1207	1509	56	613
199	1208	1512	828	965
200	1208	. 1515	3216	3812
	1 1 2 0 3	1 +2+2	1 32 10	

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
			REGION	REGION
201	1210	1516	614	790
202	1211	1522	1709	1029
203	1212	1524	614	799
204	1213	1526	3917	4081
205	1214	1529	221	2146
206	1215	1530	644	390
207	1216	1532	16	1224
208	1217	1535	885	1031
209	1218	1536	245	1156
210	1219	1538	1617	4994
211	1220	154	97	234
212	1221	1540	4325	4158
213	1222	1541	2020	2778
214	1223	1544	595	3168
215	1224	1545	328	534
216	1225	1548	47	211
217	1226	1550	49	201
218	1227	1552	418	558
219	1228	1555	509	330
220	1229	1557	699	854
221	1230	1561	847	1932
222	1231	1563	775	933
223	1232	1565	286	453
224	1233	1567	807	974
225	1234	1568	1227	1601
226	1235	1569	113	328
227	1236	157	145	2
228	1237	1570	222	845
229	1238	1572	167	685
230	1239	1574	97	1167
231	1240	1575	581	2701
232	1241	1577	1246	953
233	1242	1578	1440	175
234	1243	1579	4738	4601
235	1244	1580	1431	1568
236	1245	1581	2491	3222
237	1246	1584	463	2157
238	1247	1585	156	2366
239	1248	1586	167	691
240	1249	1587	102	305
241	1250	1589	1157	1783
242	1251	159	812	639
243	1252	1592	270	521
244	1253	1593	92	310
245	1254	1594	814	188
246	1255	1595	101	2290
247	1256	1597	119	910
248	1257	1598	178	1398
249	1258	1600	2937	2578
250	1259	1	47	
	1233	1604	4/	526

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		32, 322, 332	REGION	REGION
251	1260	1606	2204	1872
252	1261	1608	235	603
253	1262	1609	156	2366
254	1263	1611	1992	2135
255	1264	1614	968	786
256	1265	1615	2578	2751
257	1266	1616	6256	5813
258	1267	1617	29	709
259	1268	1619	1123	4071
260	1269	1621	581	2704
261	1270	1626	43	321
262	1271	1629	3616	1673
263	1272	163	509	183
264	1272	1630	81	248
265	1274	1631	9	572
266	1274	1633	2565	2807
267	1276	1634	2373	2510
268	1277	1635	3216	4508
269	1278	1636	4239	4081
270	1278	1642	4238	4020
271	1279	1643	152	304
272		1644	47	478
	1281		121	921
273	1282	1645	3815	3030
274	1283	1646	335	186
276	1284	1647	6	974
277	1285	1649	34	951
278	1286	1654	491	1387
	1287		78	560
279	1288	1656		
280	1289	1657	1431	1568
281	1290	1658	2373	1015
282	1291	1670	236	3
283	1292	1673	95	1342
284	1293	1685	2124	1786
285	1294	1690	245 977	415 774
286	1295	1691	<u> </u>	
287	1296	1699	50	247
288	1297	17	282	112
289	1298	1710	943	239
290	1299	1711	127	318
291	1300	1718	99	338
292	1301	1719	122	382
293	1302	172	33	461
294	1303	1720	180	1
295	1304	1722	160	327
296	1305	1726	175	363
297	1306	1737	84	497
298	1307	1738	188	379
299	1308	174	138	332
300	1309	1743	560	784

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
			REGION	REGION
301	1310	1747	1824	1961
302	1311	1748	97	411
303	1312	1749	151	492
304	1313	177	59	322
305	1314	1776	68	262
306	1315	1779	43	255
307	1316	178	58	399
308	1317	1781	1179	907
309	1318	1786	579	385
310	1319	1789	56	193
311	1320	180	218	78
312	1321	1800	230	394
313	1322	1801	1778	876
314	1323	181	174	428
315	1324	1829	179	42
316	1325	1846	525	785
317	1326	1848	5632	5838
318	1327	185	92	400
319	1328	1850	178	333
320	1329	186	699	1310
321	1330	1860	8	604
322	1331	1868	376	618
323	1332	187	148	366
324	1333	1870	233	388
325	1334	1872	12	206
326	1335	188	181	516
327	1336	1884	549	863
328	1337	1886	128	298
329	1338	189	28	204
330	1339	1891	11246	11097
331	1340	1895	175	417
332	1341	1897	221	400
333	1342	1899	744	890
334	1343	191	77	286
335	1344	1914	403	699
336	1345	192	8	343
337	1346	1947	656	1735
338	1347	1948	32	283
339	1348	195	129	323
340	1349	196	122	295
341	1350	1962	554	733
342	1351	1902	110	277
343	1352	1976	348	2450
344	1353	1976	93	
345	1354	1980	137	239
346				310
347	1355	2	916	13698
348	1356	20	112	303
	1357	2005	88	420
349	1358	2007	525	385
350	1359	2008	266	484

TABLE 3

SEO ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
NOCLEOTIDE	ACID	05/451,404	REGION	REGION
351	1360	2013	64	234
352	1361	2016	99	329
353	1362	2018	84	401
354	1363	2013	300	130
355	1364	2022	1240	1016
356	1365	2022	191	364
	1366	2023	231	404
357	1367	2043	3206	3349
358		2043	169	456
359	1368	2047	295	522
360	1369	<u> </u>	533	769
361	1370	2049	4	684
362	1371	205	403	699
363	1372	2051	173	379
364	1373	2055		1157
365	1374	2056	270	725
366	1375	2061	949	309
367	1376	2064	127	
368	1377	2065	248	577
369	1378	2070	204	344
370	1379	2071	374	793
371	1380	2074	945	796
372	1381	2076	300	67
373	1382	2078	416	586
374	1383	2081	316	507
375	1384	2082	20	220
376	1385	209	19	168
377	1386	210	27	395
378	1387	2102	258	452
379	1388	2104	1706	1539
380	1389	211	84	311
381	1390	212	677	231
382	1391	2120	40	414
383	1392	214	101	268
384	1393	2140	213	377
385	1394	2161	216	368
386 ·	1395	2162	106	420
387	1396	2164	104	250
388	1397	217	333	22
389	1398	218	80	325
390	1399	219	709	506
391	1400	2196	158	319
392	1401	2198	469	1164
393	1402	22	843	700
394	1403	2214	980	822
395	1404	2215	49	318
396	1405	2225	544	1974
397	1406	223	185	21
398	1407	2233	116	313
399	1408	224	189	16
400	1409	2240	2740	2525
	<u> </u>	1		

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		·	REGION	REGION
401	1410	2244	1489	1647
402	1411	2254	72	317
403	1412	226	335	120
404	1413	2260	562	738
405	1414	2268	300	67
406	1415	227	103	615
407	1416	2273	114	344
408	1417	2275	239 .	985
409	1418	2276	1358	1164
410	1419	2288	56	1459
411	1420	2291	83	532
412	1421	2296	264	530
413	1422	2298	533	781
414	1423	2300	1684	1845
415	1424	2305	8	226
416	1425	231	86	820
417	1426	232	361	1920
418	1427	233	150	467
419	1428	2331	334	2856
420	1429	2334	168	953
421	1430	2341	198	395
422	1431	2344	122	1432
423	1432	2346	1345	1187
424	1433	2348	502	729
425	1434	235	338	844
426	1435	2351	228	713
427	1436	236	232	2
428	1437	2360	1611	1357
429	1438	2362	36	263
430	1439	2364	294	1568
431	1440	2365	103	312
432	1441	2378	209	5281
433	1442	238	53	511
434	1443	2380	207	380
435	1444	239	457	663
436	1445	2392	176	2653
437	1446	2399	940	2040
438	1447	2405	144	380
439	1448	2407	1875	2702
440	1449	2415	1927	137
441	1450	242	1813	986
442	1451	2421	43	405
443	1452	2423	1556	1413
444	1453	2424	673	1041
445	1454	2432	295	1275
446	1455	2438	607	437
447	1456	2444	294	
448	1457	2444	212	437
449	1	<u></u>		1588
450	1458	2448	52	1440
	1459	2449	637	1197

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
			REGION	REGION
451	1460	245	208	876
452	1461	2450	3740	4369
453	1462	2453	222	389
454	1463	246	566	763
455	1464	2466	179	778
456	1465	2471	532	669
457	1466	2473	817	650
458	1467	2474	236	1333
459	1468	2476	173	3
460	1469	248	331	2
461	1470	2486	709	885
462	1471	249	88	456
463	1472	2496	107	1054
464	1473	2498	413	607
465	1474	2501	103	267
466	1475	2503	334	717
467	1476	2506	3740	4369
468	1477	2509	188	18
469	1478	2512	78	368
470	1479	2514	16	354
471	1480	2523	53	325
472	1481	2526	223	384
473	1482	2532	596	763
474	1483	2533	62	667
475	1484	2535	89	1519
476	1485	2537	175	375
477	1486	254	299	21
478	1487	2540	553	816
479	1488	2546	1905	1102
480	1489	2555	2046	4541
481	1490	2559	569	733
482	1491	256	9	410
483	1492	2560	288	76
484	1493	2565	3269	3502
485	1494	2569	116	478
486	1495	257	203	475
487	1496	2571	2763	2548
488	1497	2572	65	652
489	1498	2575	70	294
490	1499	2576	1195	1010
491	1500	258	434	21
492	1501	2580	155	400
493	1502	2591	53	214
494	1503	2592	163	348
495	1504	26	261	398
496	1505	2605	277	420
497	1506	261	29	598
498	1507	2614	1331	1510
498	1508	2617	235	378
	<u> </u>			458
500	1509	262	204	430

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		33,102,101	REGION	REGION
501	1510	2624	254	418
502	1511	263	247	570
503	1512	264	184	540
504	1513	2643	1108	4026
505	1514	2644	305	535
506	1515	2645	1952	1509
507	1516	2647	1225	404
508	1517	2648	41	778
509	1518	265	53	418
510	1519	2650	190	936
511	1520	2658	1576	2451
512	1521	2659	44	
513	1521	266	350	430
513	1523			153
514	1523	2663	785	1177
516	1524	2665	395	550
			41	778
517	1526	2667	244	384
518	1527	2668	174	527
519	1528	2669	27	302
520	1529	2678	1172	960
521	1530	2684	178	432
522	1531	269	341	520
523	1532	2699	1241	1083
524	1533	2701	402	2624
525	1534	2702	28	177
526	1535	2706	1108	4026
527	1536	2707	1240	1016
528	1537	271	59	346
529	1538	2714	34	987
530	1539	2715	1117	647
531	1540	2717	25	429
532	1541	2718	1670	1885
533	1542	2719	31	1137
534	1543	272	6	152
535	1544	2726	230	592
536	1545	2728	578	369
537	1546	2731	193	366
538	1547	2735	495	301
539	1548	274	352	119
540	1549	2741	94	255
541	1550	2798	1031	1240
542	1551	28	54	725
543	1552	2803	204	374
544	1553	2809	216	938
545	1554	2822	280	447
546	1555	2823	197	388
547	1556	2824	224	12
548	1557			
549	1558	2826	79	456
550	1559	2828	24	428
730	1333	2838	90	698

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
	11022	05,152,101	REGION	REGION
551	1560	284	21	197
552	1561	2847	113	262
553	1562	285	146	292
554	1563	2852	233	439
555	1564	2854	830	988
556	1565	2855	336	1043
557	1566	2856	384	614
558	1567	2857	437	748
559	1568	2859	1295	1158
560	1569	286	30	179
561	1570	2860	2618	2469
562	1571	2864	1325	1176
563	1572	2867	1034	795
564	1573	288	190	345
565	1574	2884	856	257
566	1575	2886	15	167
567	1576	2891	34	405
568	1577	2900	104	2683
569	1578	2901	193	366
570	1579	2902	91	1806
571	1580	2907	268	498
572	1581	2908	83	1564
573	1582	2910	2131	3117
574	1583	2915	715	861
575	1584	2916	52	2064
576	1585	2919	62	1015
577	1586	292	615	854
578	1587	2923	332	1279
579	1588	2924	264	422
580	1589	2925	122	1432
581	1590	2930	195	341
582	1591	2931	221	3
583	1592	2934	1642	1827
584	1593	2937	38	421
585	1594	2940	520	383
586	1595	2944	325	68
587	1596	295	49	255
588	1597	2950	226	59
589	1598	2951	110	400
590	1599	2955	303	641
591	1600	2957	365	673
592	1600	2964	96	347
593	L	2967	738	466
	1602	2968		<u> </u>
594	1603		365	428 117
595	1604	2969	<u> </u>	
596	1605	2970	314	643
597	1606	2973	961	1176
598	1607	2975	975	799
599	1608	2979	89	442
600	1609	298	152	3

TABLE 3

NUCLEOTIDE ACID 09/491,404 OF CODING REGION OF CODING REGION 601 1610 2991 112 261 602 1611 2995 201 368 603 1612 3 13559 13335 604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 <th>OP</th>	OP
REGION REG 601 1610 2991 112 261 602 1611 2995 201 368 603 1612 3 13559 13335 604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18	OTIDE
601 1610 2991 112 261 602 1611 2995 201 368 603 1612 3 13559 13335 604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617	ODING
602 1611 2995 201 368 603 1612 3 13559 13335 604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 308 88 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 <	SION
603 1612 3 13559 13335 604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619	
604 1613 30 176 751 605 1614 3002 1807 2265 606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 <t< td=""><td></td></t<>	
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606 1615 3005 339 743 607 1616 3023 64 243 608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 <td< td=""><td></td></td<>	
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608 1617 3039 71 217 609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 <t< td=""><td></td></t<>	
609 1618 304 50 334 610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 <t< td=""><td></td></t<>	
610 1619 305 226 387 611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626	
611 1620 3051 56 268 612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627	
612 1621 307 9. 278 613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628	
613 1622 308 116 274 614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629	
614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630	
614 1623 3085 97 3030 615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630	
615 1624 3088 801 634 616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631	
616 1625 3089 18 455 617 1626 3094 92 1246 618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632	
618 1627 3098 40 342 619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634	
619 1628 310 142 354 620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
620 1629 3101 48 383 621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
621 1630 3105 188 328 622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
622 1631 3107 177 413 623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
623 1632 3109 184 327 624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
624 1633 3114 70 243 625 1634 3115 295 459 626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
626 1635 3116 115 348 627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
627 1636 3119 70 222 628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
628 1637 3120 163 531 629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
629 1638 3122 60 266 630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
630 1639 3129 226 501 631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
631 1640 3146 190 363 632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
632 1641 3151 212 1588 633 1642 3153 86 517 634 1643 3165 244 453	
633 1642 3153 86 517 634 1643 3165 244 453	
634 1643 3165 244 453	
625	
635 1644 317 97 342	
636 1645 3179 106 873	
637 1646 3181 108 896	
638 1647 3182 554 775	
639 1648 3192 268 441	
640 1649 3194 923 1192	
641 1650 3195 38 376	
642 1651 32 185 334	
643 1652 3200 199 561	
644 1653 3201 516 848	
645 1654 3202 232 681	
646 1655 3208 836 633	
647 1656 3210 202 384	
648 1657 3214 349 588	
649 1658 3215 859 380	
650 1659 3216 51 320	

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
MOCHEOTIDE	ACID	05/451,404	REGION	REGION
651	1660	3220	116	283
652	1661	3222	324	545
653	1662	3227	385	1197
654	1663	323	65	223
655	1664	3240	385	1197
656	1665	3243	65	916
657	1666	3250	263	463
658	1667	3250	244	480
659	1668	3252	136	297
660	1669	3254		439
	<u> </u>		83	
661	1670	3255	573	920
662	1671	3257	548	757
663	1672	3259	34	822
664	1673	326	58	525
665	1674	3263	102	350
666	1675	3270	313	152
667	1676	3271	117	473
668	1677	3272	44	190
669	1678	3273	106	486
670	1679	3274	246	392
671	1680	3278	174	1
672	1681	3281	988	1134
673	1682	3282	101	334
674	1683	3291	129	284
675	1684	3294	101	595
676	1685	3296	107	565
677	1686	3298	130	552
678	1687	3299	333	515
679	1688	3300	324	121
680	1689	3303	378	157
681	1690	3306	296	637
682	1691	3307	1454	1660
683	1692	3309	163	471
684	1693	3311	335	478
685	1694	3312	5	280
686	1695	3313	298	546
687	1696	3314	50	526
688	1697	3315	99	413
689	1698	3322	101	685
690	1699	3323	66	356
691	1700	3324	76	462
692	1701	3328	248	904
693	1702	3335	136	393
694	1703	3336	47	733
695	1704	3338	181	786
696	1705	3339	58	231
697	1706	3342	226	390
698	1707	3342	72	488
699	1708	3356	208	384
700	1709		194	436
	1 1 / 0 3	3358	1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	1 430

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
			REGION	REGION
701	1710	3360	263	1459
702	1711	3366	55	816
703	1712	3367	364	735
704	1713	3370	237	878
705	1714	3371	188	721
706	1715	3372	14	241
707	1716	3373	42	290
708	1717	3387	32	202
709	1718	3389	29	256
710	1719	3390	181	393
711	1720	3396	520	822
712	1721	3410	10	153
713	1722	3412	82	291
714	1723	3414	453	292
715	1724	3421	158	337
716	1725	3427	430	618
717	1726	3430	210	380
718	1727	3431	295	432
719	1728	3440	419	556
720	1729	3444	402	256
721	1730	3445	281	430
722	1731	346	42	722
723	1732	347	384	689
724	1733	3470	114	530
725	1734	3478	38	217
726	1735	3479	161	379
727	1736	348	37	231
728	1737	3482	156	296
729	1738	35	255	575
730	1739	3503	185	454
731	1740	3505	252	422
732	1741	3529	37	183
733	1742	353	262	522
734	1743	3537	127	273
735	1744	3539	98	268
736	1745	3542	25	312
737	1746	3543	70	228
738	1747	3544	31	177
739	1748	3548	972	385
740	1749	3553	27	164
741	1750	3560	113	358
742	1751	3563	483	764
743	1752	3564	6	434
744	1753	d		L
745	1754	3566	316	507
746	1755	3570	6	377
747	1756	3574	108	440
748	<u> </u>	3576	569	348
	1757	3579	293	442
749	1758	3582	20	388
750	1759	3583	172	396

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		05, 152, 151	REGION	REGION
751	1760	3587	84	449
752	1761	3596	91	459
753	1762	3599	40	474
754	1763	3606	335	1105
755	1764	3609	169	666
756	1765	3617	141	410
757	1766	3620	218	388
758	1767	3630	189	1
759	1768	3642	122	643
760	1769	3644	431	664
761	1770	3647	274	720
762	1771	3651	245	472
763	1772	3652	259	642
764	1773	3653	153	1994
765	1774	3654	87	554
766	1775	3654	57	2744
767	1776	3658	387	920
768	1777	366	402	578
769	1778	3660	120	530
770	1779	3661	480	674
·			1096	938
771	1780	3663	<u> </u>	1015
772		3669	689	
773	1782	3677	469	642 889
774	1783	3678	1194	1134
775	1784	3685	233	706
776	1785	3689		.l
777	1786	3693	21 55	446
778	1787	3699		
779	1788	370	59	262
780	1789	3707	38	436
781	1790	3711	229	474
782	1791	3713	314	463
783	1792	3717	178	675
784	1793	3720	258	695
785	1794	3721	96	548
786	1795	3722	32	562
787	1796	3724	220	513
788	1797	3726	180	467
789	1798	3729	251	523
790	1799	373	110	340
791	1800	3735	91	636
792	1801	3736	275	880
793	1802	3738	106	621
794	1803	3762	702	1175
795	1804	3768	293	598
796	1805	377	96	257
	1 7003			
797	1806	3772	169	2
L		3772 3786	169	2 248
797	1806			

TABLE 3

OF NUCLEOTIDE OF AMINO ACID IN USSN 09/491,404 NUCLEOTIDE OF CODING REGION REGION REGION NUCLEOTIDE OF CODING REGION REGION REGION REGION NUCLEOTIDE OF CODING REGION REGION REGION REGION NUCLEOTIDE OF CODING REGION REGION REGION REGION 801 1810 379 248 421 421 802 1811 38 146 3 146 3 140 34 275 3	SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
NUCLEOTIDE ACID 09/491,404 OF CODING REGION OF CODING REGION 801 1810 379 248 421 802 1811 38 146 3 803 1812 382 24 275 804 1813 385 138 1 805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 813 1824 416 736 909		1		1	i
REGION R	NUCLEOTIDE	1		1	
801 1810 379 248 421 802 1811 38 146 3 803 1812 382 24 275 804 1813 385 138 1 805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824			, , , , , , , , , , , , , , , , , , , ,		i e
802 1811 38 146 3 803 1812 382 24 275 804 1813 385 138 1 805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 813 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827	801	1810	379		
803 1812 382 24 275 804 1913 385 138 1 805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 819 1828	802	1811		146	1
804 1813 385 138 1 805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 819 1828 435 140 445 820 1829	803			1	
805 1814 388 268 74 806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 819 1828 435 140 445 820 1829 437 160 423 821 1831	804				
806 1815 39 302 3 807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 817 1828 435 140 445 820 1829 437 160 423 821 1830 439 347 706 822 1831 <td>805</td> <td></td> <td>.4</td> <td>1</td> <td>L</td>	805		.4	1	L
807 1816 391 24 368 808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 819 1828 435 140 445 820 1829 437 160 423 821 1830 439 347 706 822 1831 44 91 282 823 1833 <td>L</td> <td></td> <td>-l</td> <td></td> <td></td>	L		-l		
808 1817 395 51 482 809 1818 397 422 766 810 1819 399 102 311 811 1820 4 11219 13123 812 1821 405 253 2 813 1822 406 342 665 814 1823 411 321 542 815 1824 416 736 909 816 1825 422 1541 867 817 1826 43 330 686 818 1827 434 207 34 819 1828 435 140 445 820 1829 437 160 423 821 1830 439 347 706 822 1831 44 91 282 823 1832 450 136 402 822 1831 </td <td></td> <td></td> <td></td> <td></td> <td></td>					
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TABLE 3

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855	1864	540	270	1
856	1865	541	38	412
857	1866	546	388	2
858	1867	555	199	438
859	1868	556	144	482
860	1869	559	380	165
861	1870	563	27	617
862	1871	566	158	382
863	1872	568	69	320
864	1873	57	6	158
865	1874	571	18	1516
866	1875	572	32	505
867	1876	573	139	456
868	1877	574	49	771
869	1878	576	519	370
870	1879	578	168	1
871	1880	580	159	641
872	1881	581	108	497
873	1882	582	80	403
874	1883	587	172	435
875	1884	589	27	374
876	1885	590	84	428
877	1886	595	68	1138
878	1887	598	1023	766
879	1888	61	65	208
880	1889	612	310	546
881	1890	614	166	918
882	1891	617	252	602
883	1892	62	969	661
884	1893	620	188	418
885	1894	622	877	1014
886	1895	629	202	687
887	1896	63	98	277
888	<u> </u>	632	221	367
889	1897 1898	64	536	381
890		640	338	3
891	1899	641	12	395
L	1900		194	395
892	1901	642		<u> </u>
893	1902	644	15	395
894	1903	646	132	380
895	1904	647	3	389
896	1905	650	135	413
897	1906	651	231	428
898	1907	653	128	442
899	1908	654	214	77
900	1909	656	49	465

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
		02,122,101	REGION	REGION
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902	1911	66	267	614
903	1912	662	387	701
904	1913	666	76	498
905	1914	667	517	2184
906	1915	668	1423	788
907	1916	67	107	622
908	1917	678	172	387
909	1918	68	78	341
910	1919	680	832	671
911	1920	683	505	164
912	1921	687	105	521
913	1922	690	139	294
914	1923	691	244	456
915	1923	699	194	754
916	1925	701	371	520
917	1926	702	1888	2028
918	1927	704	1254	808
919	1928	704	126	1463
920	1929	706	31	
921	1930	707		390
922			1152	2
923	1931	709		934
924	1932	716	744	541
925	1934	722	1360	1220
926	1934	725	173	430 271
927	1936	727	498	
928	1936		18	164
929	1937	729	230	3
930	<u> </u>	.i	262	834
931	1939	731	491	246
	1940	740	20	322
932 933	1941	741	1430	1167
933	1942 1943	747	660	523
935		749	263	727
936	1944	750	209	391
	1945	751	753	517
937	1946	755	172	387
938	1947	756	209	376
939	1948	76	656	513
940	1949	760	131	538
941	1950	763	893	1126
942	1951	766	1271	1537
943	1952	771	458	318
944	1953	775	391	558
945	1954	781	410	1684
946	1955	791	967	1284
947	1956	793	554	970
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949	1958	796	342	199
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TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
OF	OF AMINO	IN USSN	NUCLEOTIDE	NUCLEOTIDE
NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
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955	1964	808	210	905
956	1965	812	162	920
957	1966	819	723	2669
958	1967	820	964	725
959	1968	825	182	328
960	1969	829	1843	2292
961	1970	830	58	201
962	1971	832	150	341
963	1972	835	130	762
964	1973	836	449	291
965	1974	838	175	324
966	1975	84	175	435
967	1976	842	73	393
968	1977	844	423	824
969	1978	845	214	32
970	1979	846	120	317
971	1980	847	212	364
972	1981	85	190	426
973	1982	852	74	541
974	1983	855	1653	1465
975	1984	857	1964	2659
976	1985	858	598	1020
977	1986	861	58	933
978	1987	876	222	779
979	1988	878	2021	2161
980	1989	879	189	362
981	1990	88	39	278
982	1991	886	1165	1022
983	1992	891	158	310
984	1993	892	759	995
985	1994	895	224	379
986	1995	897	131	622
987	1996	9	1678	1448
988	1997	901	55	753
989	1998	906	450	623
990	1999	913	40	237
991	2000	918	17	334
992	2001	92	385	122
993	2002	926	772	518
994	2002	929	146	283
995	2004	932	23	175
996	2005	934	38	235
997	2006	935	286	423
998	2007	936	24	284
999	2008	939	450	623
1000	2009	94	139	2
	12009	1 24	133	14

TABLE 3

SEQ ID NO:	SEQ ID NO:	SEQ ID NO:	START	STOP
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NUCLEOTIDE	ACID	09/491,404	OF CODING	OF CODING
			REGION	REGION
1001	2010	944	156	860
1002	2011	947	174	356
1003	2012	957	80	400
1004	2013	96	187	387
1005	2014	964	1352	1528
1006	2015	97	166	2
1007	2016	98	535	344
1008	2017	995	559	386
1009	2018	997	34	231

WHAT IS CLAIMED IS:

1. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of SEQ ID NO: 1-1009, a mature protein coding portion of SEQ ID NO: 1-1009, an active domain of SEQ ID NO: 1-1009, and complementary sequences thereof.

- 2. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide hybridizes to the polynucleotide of claim 1 under stringent hybridization conditions.
- 3. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide has greater than about 90% sequence identity with the polynucleotide of claim 1.
- 4. The polynucleotide of claim 1 wherein said polynucleotide is DNA.
- 5. An isolated polynucleotide of claim 1 wherein said polynucleotide comprises the complementary sequences.
- 6. A vector comprising the polynucleotide of claim 1.
- 7. An expression vector comprising the polynucleotide of claim 1.
- 8. A host cell genetically engineered to comprise the polynucleotide of claim 1.
- 9. A host cell genetically engineered to comprise the polynucleotide of claim 1 operatively associated with a regulatory sequence that modulates expression of the polynucleotide in the host cell.
- 10. An isolated polypeptide, wherein the polypeptide is selected from the group consisting of:
 - (a) a polypeptide encoded by any one of the polynucleotides of claim 1; and

(b) a polypeptide encoded by a polynucleotide hybridizing under stringent conditions with any one of SEQ ID NO:1-1009.

- 11. A composition comprising the polypeptide of claim 10 and a carrier.
- 12. An antibody directed against the polypeptide of claim 10.
- 13. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polynucleotide of claim 1 for a period sufficient to form the complex;
 and
- b) detecting the complex, so that if a complex is detected, the polynucleotide of claim 1 is detected.
- 14. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample under stringent hybridization conditions
 with nucleic acid primers that anneal to the polynucleotide of claim 1 under such
 conditions;
- b) amplifying a product comprising at least a portion of the polynucleotide of claim 1; and
- c) detecting said product and thereby the polynucleotide of claim 1 in the sample.
- 15. The method of claim 14, wherein the polynucleotide is an RNA molecule and the method further comprises reverse transcribing an annealed RNA molecule into a cDNA polynucleotide.
- 16. A method for detecting the polypeptide of claim 10 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex; and

b) detecting formation of the complex, so that if a complex formation is detected, the polypeptide of claim 10 is detected.

- 17. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10 under conditions sufficient to form a polypeptide/compound complex; and
- b) detecting the complex, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 18. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10, in a cell, under conditions sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and
- b) detecting the complex by detecting reporter gene sequence expression, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 19. A method of producing the polypeptide of claim 10, comprising,
- a) culturing a host cell comprising a polynucleotide sequence selected from the group consisting of a polynucleotide sequence of SEQ ID NO: 1-1009, a mature protein coding portion of SEQ ID NO: 1-1009, an active domain of SEQ ID NO: 1-1009, complementary sequences thereof and a polynucleotide sequence hybridizing under stringent conditions to SEQ ID NO: 1-1009, under conditions sufficient to express the polypeptide in said cell; and
 - b) isolating the polypeptide from the cell culture or cells of step (a).
- 20. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 1010-2018, the mature protein portion thereof, or the active domain thereof.

21. The polypeptide of claim 20 wherein the polypeptide is provided on a polypeptide array.

- 22. A collection of polynucleotides, wherein the collection comprises the sequence information of at least one of SEQ ID NO: 1-1009.
- 23. The collection of claim 22, wherein the collection is provided on a nucleic acid array.
- 24. The collection of claim 23, wherein the array detects full-matches to any one of the polynucleotides in the collection.
- 25. The collection of claim 23, wherein the array detects mismatches to any one of the polynucleotides in the collection.
- 26. The collection of claim 22, wherein the collection is provided in a computer-readable format.
- 27. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.
- 28. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising an antibody that specifically binds to a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.

SEQUENCE LISTING

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<213> Homo sapiens

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<212> DNA

<213> Homo sapiens

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	acattgacga					960
	ttcgttcttc					1020
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1740

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780

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gagtaccgaa accactggtc ggacgtgctg gctggcttcc tgacaggggc ggccatcgcc
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                                                                      420
                                                                      480
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tacccetgce agegeecace eccagecagg geceetegce tteeteecet ggacetgggg
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                                                                      780
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                                                                      840
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                                                                     1320
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                                                                     1380
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                                                                     1560
cctgttcttc acactgcccc ccatgcctca gcctcataca gatgtgccat catggggggc
                                                                     1620
atgggtggag caaaggggct ccctcacccc gggcaggcaa aggcagtggg tagaggaggc
actgecece ttteetgece ectecteate tttaataaag acetggette teatetttaa
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                                                                     1712
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<210> 46 <211> 755 <212> DNA <213> Homo sapiens

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120 ggaggggctg qccaggggtt ggtcagagga cactgaaggt ctcagagcct gctccattac 180 gggtqqqcaq aqcccttcct caaqccttgt taggagccag acctcactgt gtattcccag 240 gaggggaggt teteggagte gaggeageat ttggateeag tttcattete ageacettet 300 tectacacca gecattatte ttteetggee ecaaacteag ggeaacecaa tatttgatat 360 catctgaccc cactcacttg ccagctggac ggggccccaa cagtgtctcc atgtaaagga 420 tgcagctttc caatcccacc caatctttgt gcacctactg tgtgctggcg ctggaagcag 480 ggaqcaggag aggatgactc agttctttat cacagataat gggcacagct catatttatc 540 600 gecagettea tttateetgg gtaetgagaa cattgtaatg cacettteae cetteaegge 660

60

720

755

gecagettea tttatectgg gtaetgagaa cattgtaatg cacettteae eetteaege gtattgtget ttgaegeegg aacttttggga agecaaggag gaetattace ttateteaga tgggggaeca gteeggaeaa tegaaggtee tettttettg gtaeeggeae attgttaeee gattgggegg eeegetggtt ateetttaat acaae

<210> 47 <211> 2820 <212> DNA <213> Homo sapiens

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gcaactgagg gcccatttgc tatggatcca gattctggct tcctgctggt gaccagggcc
                                                                      300
ctggaccgag aggagcaggc agagtaccag ctacaggtca ccctggagat gcaggatgga
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                                                                      420
ccccatttct ctcaagccat ctacagagct cggctgagcc ggggtaccag gcctggcatc
                                                                      480
cccttcctct tccttgaggc ttcagaccgg gatgagccag gcacagccaa ctcggatctt
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cgattccaca tcctgagcca ggctccagcc cagccttccc cagacatgtt ccagctggag
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                                                                      720
caggccactg ccaccgtgga agtctccatc atagagagca cctgggtgtc cctagagcct
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cagagectgg acaccaactt tatggactge ceatgggagt getecaaatg teagggtgtt
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<210> 48
<211> 1517
<212> DNA
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<213> Homo sapiens

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<400> 48

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ttgtattatg acacatatgc acaaggatta gctctatagc gcgctgtaca tggtgggtcc 180
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agettgetee ceagtagttg tttgagteea gattetttgg ggtggateet etttteagag
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ctcctccaac attagcataa ttaaagccaa ggaggaggag gggggtgagg tgaaagatga
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gctggaggac cgcaataggg gtaggtcccc tgtggaaaaa gggtcagagg ccaaaggatg
                                                                      420
ggaggggtc aggctggaac tgaggagcag gtgggggcac ttctccctct aacactctcc
                                                                      480
cctgttgaag ctctttgtga cgggcgagct caggccctga tgggtgactt cgcaggcgta
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gactttgtgt ttctcgtagt ctgctttgct cagcgtcagg gtgctgctga ggctgtaggt
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                                                                      660
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                                                                     1200
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                                                                    1440
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tgagctgggg cttccat
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<210> 49 <211> 1614 <212> DNA

<213> Homo sapiens

<400> 49

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cacceaaage tecaetgttg aeggacgggg aaaagecaga aeegaceget etet
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     <211> 659
     <212> DNA
     <213> Homo sapiens
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ggcagccctg ggctgctgtg aggtcctgag cacagtgcat cctgaggaga cagtgctgcq
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ggccccgcct actaacttcc agagatgtca gctgcagcag ggcagcgccc tqqttaqaqa
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gacggcatgg ggagttggca gggggaggcc ctcggagaga tggcatgggg agttggcagg
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                                                                      360
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                                                                      420
ttgaacaaag actttgtttg tgccttcatt cgttcagcac atgtttacaq tqtqcctqtq
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atgtcccagg cgcactgccc tattcttgac atccttgtgg tgggatcaac tgcttgcctg
                                                                      540
tocatagogo aggocattac tagaggtgtt ttotgggggg cgaacacogt tottttgcag
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     <210> 51
     <211> 450
     <212> DNA
     <213> Homo sapiens
     <400> 51
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acacggacag accgctgtag aagaggcttt ggtgtctgta ctgctcagaa gggcgaggca
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tgcatgctct taaggattta ccagegcaat actetecaga tateatacat ggtgtgtcag
                                                                      240
aaattetgea gagacatgae atttgatete aggaategga ettatgttea tacatgetge
                                                                      300
aactacaatt actgtaactt taaactctaa gatatttgcc ctcctgaggt ctcgctttgg
                                                                      360
aatgtcccca atgttgctca tccttcacac tctgctggcc cttgcttccc ttccgtgtct
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gtcctgacaa tacccctgcc ctcgcattaa
                                                                      450
     <210> 52
     <211> 1044
     <212> DNA
     <213> Homo sapiens
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     <221> misc_feature
     <222> (1)...(1044)
     <223> n = a,t,c or g
     <400> 52
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ttctccctct gtattatctc agcatacact gagettgcaa acatatgaat ttcacattgt
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egtggaatet tacageetge taetteetaa gttttettta gacaagetge ettggtgace
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300
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aatttgtget tttaaaagta tettacaett tteaettatt tgtaeettta aaaaaatett
                                                                      420
ttttttttt taaaccaaag gtttgcagta tcttcaaagt ctgaattttg ageggatagg
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gatgagecae ctaaateece tgaaaatttg cetgeeetea ggggttaaet tttttgetge
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aatcacaaag taggttattt acgctttctt gatgggagtt attaaaaaaa ttttaattta
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conceptigite tetactetet gitteegeact caegetteat acatteetag aegeeegege
                                                                      960
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ctacctcacg caccgactcc acca
                                                                     1044
```

<210> 53
<211> 1328
<212> DNA
<213> Homo sapiens
<220>
<221> misc_feature
<222> (1) ... (1328)
<223> n = a,t,c or g

<400> 53

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<210> 54 <211> 804 <212> DNA <213> Homo sapiens

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gaggcagetg ccgagetggc ccaggtgctg ccgcagatgg gccggctgaa gagagtggag
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     <212> DNA
     <213> Homo sapiens
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<400> 67

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gctggtggag cctggggtta cagcattggg aagaaatgga gatggagaac aggacagctg
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taaataccat aacctgtccc tcccaccccc caactacatt cgaaaaagta agaacagcag
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                                                                      480
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ctagggatca aaactcctta gtctggttga ttgctgaatg ggagaggagt aagtgagaaa
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gatcatggca ggctggccct gcaattattc aaacccaggc ccctggctgc ctgggaacgg
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gcaaagcagg ccaaacttag cttccatggt tacatttgga agtttctatt catgacacca
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aataaaagtg gggaagaagg aagcatggct tactgaagta gtctcaggaa gacagggcaa
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gctqcatact ctttatttat qttaaaacaa qtaqaaccca ccaaattaat tacaaqataq
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aacagaaaca gattaaaata catcagctgg tttgtgttta gaagaggtaa tgagacaact
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attacagtaa gatattttga attaagaaac aaggtgtaaa ctgtaggaaa atatacaaat
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aaacacaact gaaataaaaa aaaaaaaaa
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<210> 68
<211> 533
<212> DNA
<213> Homo sapiens
```

<400> 68

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tgatcctgtg gcttcagttg agctgggttt ggagccaaca gaaggaggtg gagcagaatt 180

ctggacccct cagtgttcca gagggagcca ttgcctctct caactgcact tacagtgacc 240

gaggttccca gtccttcttc tggtacagac aatattctgg gaaaagccct gagttgataa 300

tgtccatata ctccaatggt gacaaagaag atggaaggtt tacagcacag ctcaataaag 360

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420
ccagccagta tgtttctctg ctcatcagag actcccagcc cagtgattca gccacctacc
tetgtgeega ttatteagga aacacete ttgtetttgg aaagggeaca agaetttetg
                                                                      480
                                                                      533
tgattgcaaa tatccagaac cctgaccctg ccctgtacca gctgagagac tct
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     <211> 850
     <212> DNA
     <213> Homo sapiens
     <400> 69
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ctaaccacct gttcctattt tcctaccctt cattaatttg acttttgact tttgataaag
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                                                                      240
ttatcacata ttaaaatata cgtgggtgct aagcettata ctgtgaatgt tccagggttc
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aaccttggga gtccaaagct catcctaaga ggaattaata atatatcttt ttttttttgg
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gcccaggcgg gggggctaag gcctgaaacc ccagcacttg ggaagcccaa ggcaggggga
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                                                                      600
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                                                                      720
ccttggactc aagccggggg agacgaacgg gacccctccc aaaaaaaaa aagggggggc
                                                                      780
ccttaagggg aaccattgta ccgcggcggc ggggggatga gccttttaag ggcaccaaac
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cccgggcggc
                                                                      850
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     <211> 859
     <212> DNA
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     <223> n = a,t,c or g
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                                                                      300
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ataaagcact gatgaaagat ccaaggtttt ggatagcaat tgctgcatat ttagcttgtg
                                                                      360
                                                                      420
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taagtcaatt aataaaaaga gccatctgga ggaaataaaa aaaaaaggaa gactctatga
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                                                                      660
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ctccgcgcgg gtggggaaaa ataatttttt tattggggcc caaaaataaa ttcccgggcc
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                                                                      780
egececteg teegeeggeg teecegeteg geggeeteeg geceegeggt eeegegggee
                                                                      840
                                                                      859
cggccccggc gggtagccg
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<210> 71
     <211> 864
     <212> DNA
     <213> Homo sapiens
     <400> 71
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                                                                      120
caggtgctga tccttttaga ggcagccgtg gaagaggaaa agagatcaga agagaaaagg
                                                                      180
atattggttt getggaacat gtgggacaag aagtteecag aagaatttgt gageaactte
                                                                      240
ccgacagtaa ggccctggct agacctcagg atggtccctg cctcctggac attaggaagc
                                                                      300
ccaaaggcca qaacaaaaac acatgcctag tqqqqqaaqq ctcactaaqa qqqcaccaaq
                                                                      360
tggggcaaat acccetggta acccatttat ggaggetgee acagaaatge tagttggaaa
                                                                      420
ttttcctcct tcaqtctatc atqaatttct tttttctctt ttqaqatqaa qtcqccqqq
                                                                      480
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cctqaccttt qqaqqaacca cccatcttqq cctccaqacq qqctqcqatq qaaqcttqaq
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ccactgtagc tcgatgtacc gtgaatatta gctttagggc agttttaagt gggggagact
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ttaacaggac agtttacacg tataatccca aacaccccc gggctgcgcc tgqtqqaqaq
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gaaaatgtat tgattatgaa aacc
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     <210> 72
     <211> 746
     <212> DNA
     <213> Homo sapiens
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aacatcatca gcccaagcaa caatggtggc aatgttcagg agacagtgac aattgataat
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gaaaaaaata ccgccatcat taacatccat gcaggatcat gctcttctac cacaattttt
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gactataaac atggctacat tgcatccagg gtgctctccc gaagagcctg ctttatcctg
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aagatggacc atcagaacat ccctcctctg aacaatctcc aatggtacat ctatgagaaa
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caggetetgg acaacatgtt etecageaaa tacacetggg teaagtacaa eeetetggag
                                                                      420
tctctgatca aagacgtgga ttggttcctg cttgggtcac ccattgagaa actctgcaaa
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catatccctt tgtataaggg ggaagtggtt gaaaacacac ataatgtcgg tgctggaggc
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tgtgcaaagg ctgggctcct gggcatcttg ggaatttcaa tctgtgcaga cattcatgtt
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taggatgatt agecetettg ttttatettt teaaagaaat acateettgg tttacaetea
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aaagtcaaat taaattottt cocaatgooc caactaattt tgagattcag tcagaaaata
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taaatqctqt atttataaaa aaaaaa
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     <210> 73
     <211> 1928
     <212> DNA
     <213> Homo sapiens
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     <221> misc feature
     <222> (1)...(1928)
     \langle 223 \rangle n = a,t,c or g
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gaatgactca taaatcaatg caggagcagt tagcagacca cggctgtatg gctcagtgtt
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tttaagagtg aaagagaaaa ttctatttta actaaaacta aggcttaatt tttaaatcca
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cagaggtacc aaggcgccct ctaatggtga actcaaacaa tgctctattt tgtaatgagc
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tacagtttca gttagaaatt gtggtaaatt cgttagggaa ttatgaacag attttttct
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ttttttgtaa aggetttata atttettaat ggttggeeat eagttttgte tettetatge
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attttcaggc tgtattctac aaggettctt gcctattggt gaagggttat tgggggtttg
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tctgtaatgg ttattgcact gattattttt cttaggtccc cagccatggc tgggggatta
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aagaacctgc acagatttac tcatattcct tcaggaaagt gtttagatcg ctcagaggtc
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atgaaatc
                                                                     1928
```

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<210> 74
<211> 3644
<212> DNA
<213> Homo sapiens
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ggaagattgc ttcagcccag gagttcgaga ccagcctggg caacacagga agaccccgtc
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711

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420

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780

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	ataaagcaag					7620
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	acccaggaat					7740
	ccacccccaa					7800
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	ccatgctcat					9660
	tacagattca					9720
	actactttaa					9780
	aagttcatat					9840
	aaagctggag					9900
	acagcatggt					9960
	gaaataatac					10020
	tggggaaagg					10080
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agatgctgga tcctccaggc tggggtagaa ttgcaacagc ttgtccttcc ttgtgggtgc
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                                                                      240
catgtccgcc aggggtcctg gccatgcctg cccgaccaag gagtaggtcc gggaccccgt
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caactaggaa cagaccctgg ccaggagcgg tggctcacgc ctataatccc agcacgttgg
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     <213> Homo sapiens
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gttcaccage tttegggeta getgggtagg aggtgatget geceeggtet ggeacceaet
                                                                      300
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tacctggett tecetaatte accgaeceea ggattaacee catggtggtt ggtateaggg
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                                                                      420
teceggtace atetytycee ageeggggae tyggaacety gtttetecat gaggageeat
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cccagggcct gcaggaggga ctagaagcca gaggactctg aggctccgct tcctggggac
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actgatectg tgtggtgtte accceaggae gtgggagget getetgteee tetggeetta
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                                                                     1200
                                                                     1260
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1500

1560

1620

1667

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cgtagcggaa tcgggggtat gctgttcgaa ttcataagaa cagggaggtt agaagtaggg
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tcttggtgac aaaatatgtt gtgtagagtt caggggagag tgcgtcatat gttgttccta
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ggaagattgt agtggtgagg gtgtttatta taataatgtt tgtgtattcg gctatgaaga
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ataaggcgaa ggggcctgcg gcgtattcga tgttgaagcc tgagactagt tcggactccc
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cttcggcaag gtcgaagggg gttcggttgg tctctgctag tgtggagata aatcatatta
                                                                    420
480
                                                                    540
gagaggttaa aggagccacc ttattagtaa tgttgatagt agaatgatgg ctagggtgac
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cttcatatga gattgtttgg ggctacctgc tccgcagtgc gccgatcagg gcgtagtttg
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tggccatggg tatgttgtta agaagaggaa ttgaacctct gactgtaaag ttttaagttt
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     <211> 849
     <212> DNA
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     <221> misc feature
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     <223> n = a,t,c or g
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ctatctgctg tgcctcagct gcttatttgg gacaggtatg gttacttata tatgcctggc.
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ccagagcatt ttaaattaat cctttctgtt tcattattcc tcacttacac ttaaaatgac
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cccgccqgcg cggacgcgag ctctgtcgca ccgatagaac cgacgcatgg cgccgataca
                                                                    780
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cgacggacg
                                                                    849
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    <211> 860
    <212> DNA
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atcccatttg tgggttctct taattctatc attgcttctt ttcctgcgga aaagttttaa
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gttttatgca gtctcatttg tgtgttttgc ttttgttgcc ttttggaata atctacagaa
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aatcataget caggecaatg teatacagte teettetata ttteettgta gtagttetae
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ttcttctgtg cccaaaaaca ttattgaaca agaccaagaa cacttaaaac ggaaacaaat
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tgggcaagtc ttaagcaagc cattcctgct ttctgggcct ggctcccatg ggccattaga
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ccggatccat gggcccagca cccaggggca tcagcttctg cttttatggg tgggggtctt
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gttcacctgg gcaaaggcat cgcttaagtg ctgaccttta atcagatcca cgatgcgcag
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gcaataaggt aagaaaataa aaatacaaaa atcaacatac aaccaactgc aaaggaaatt
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ttaaaaaatt acattcacaa atagcataaa aagaataaag gatttagaaa taaagttaat
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taaataaatg gagatatgtc ccatgtttgc aaataggaaa atacagtatc atcaaggtgt
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360

420

480

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<211> 768

<212> DNA

<213> Homo sapiens

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<212> DNA
<213> Homo sapiens
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<212> DNA

<213> Homo sapiens

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<223> n = a,t,c or q
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                                                                    540
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1140
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                                                                   1440
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                                                                   1620
accatacttg aageteecag gacaagggtt gagaggetea acceetettt cagettetat
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     <213> Homo sapiens
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<210> 182

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<210> 183 <211> 1298 <212> DNA <213> Homo sapiens

<400> 183

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1020
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cccacagett tgatggcagt gcctcatett caacttttgt gctcccettt gcctaaacce
tatggcctcc tgtgcatctg tactcaccct gtaccacaaa cacattacat tattaaatgt
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ttctcaaaga tggaaaaaaa aaaaaaaggg gggccccttt taagggacca agttttacta
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    <211> 797
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    <213> Homo sapiens
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teteceaete teagtggett etteaeggta tgttetgtet eteaeettea egtteeeegg
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                                                                    660
cactegeeee tgaegggaae gagaeeeagg gaettetgee eeaeeaggea teeteggtgt
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                                                                    180
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cattctgctg tgcagaccac cttttaaaag tgatcacaaa ccatttgctg aatacttgtg
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660
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aggcagatca cgaagtcagg agattgagac catcctggct aacatggtga aaccccatct
                                                                    720
                                                                    780
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                                                                     1260
                                                                     1320
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aaggtgteet tgattttggt tageetggte ttttetetgt gatetetete atgagttett
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                                                                     1560
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atatattaag taaaaggaaa tgagtottga gaacattgag aaatggaaac gtttgagtag
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gcccaggtgc ggggggctca tgtctggaaa tccccatcat ggtgggaggg cccagcgtgg
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     <221> misc_feature
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                                                                       60
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                                                                      120
cccctgagcc tggtccttct cgtgtgtggc tggatctgcg gcctgctcag ctccctggcc
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gtgtcagttt catgtaccag cccgaggaca gcagcaatga aggggaaggc gtgggagagg
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ccggggagga agacggtaaa gacaccaggg aggaggacgg tgcggacgcc agcgaggaag
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                                                                 1020
ccagcgagga agatgataca attaccaatg aaaaggcaca cagtattcta aattttttga
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agtcaacaca ggaaatgtat ttttatgacg gtgtctccag agatgcagct tcagctgccc
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<213> Homo sapiens
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<210> 202
<211> 3878
<212> DNA
<213> Homo sapiens
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<400> 202

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<213> Homo sapiens

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                                                                      540
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                                                                      120
accaceccaa gggateetee ctateacett teeecetget etteeteeta etttgtaaaa
                                                                      180
gagggetttt etgtggttta geacttgaat ttetgeagta egttgattet gaegeteata
                                                                      240
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                                                                      300
aatcacctca aatacaacqt aatqttqqqt ctaataaqqa aactccactc tqctccactt
                                                                      360
taggaagaaa tegttgetag gaacaacaca tattaaactg etetatgeta tttatcagat
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tcaggagttc aagaccagcc tggccaatat ggtgaaatcc cgtctctacc gaaaatacca
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gagaatcact tgaacccagg
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caggegtggg ggtgcatgcc tgtaatccca gctaataaaa aggctgaggc aggagaatca
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cgattgggtg gttggagcca ggactgctgg ggaggaggcg gctgcagcca gcagctgaca
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                                                                      660
teteaquetq etgeaggace tgteteacte cagaaageat gageeeteec cacetggagg
                                                                      720
ctgcacaggt aagcetetga aateccaagg cataaagtee catggaagee getteetetg
                                                                      780
caaggccaaa tacatacgtc acagaaccca ataaggtcct acagcaaatt cgacaggcct
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1440

1500

1560

1620

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<400> 221

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caggetggge aatatttgga gacegggggt geagecette tgtgggeegt teeceetttg
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                                                                     1980
                                                                     2040
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gcactttagg aggccaccac aggaggatca ctcccgtgat caaaaccaac ctgggcaaca
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<210> 222 <211> 1947 <212> DNA <213> Homo sapiens

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<210> 223 <211> 1131 <212> DNA

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1947

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coefficient coefficient theoretical taaaaattat aatotottaa tittottigaa
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<213> Homo sapiens

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<211> 923
<212> DNA
<213> Homo sapiens
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                                                                      480
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                                                                      540
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                                                                      923
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<210> 274

<211> 4784

<212> DNA

<213> Homo sapiens

<220>

<221> misc_feature

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<223> n = a,t,c or g
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<212> DNA
<213> Homo sapiens
<220>
<221> misc_feature
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<223> n = a,t,c or g
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<213> Homo sapiens

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tgtcttctaa cagagtatgt acaggaaggt aatagttgct ttaacagtgt tcaqacttca
                                                                      300
aaagtgtagc tgttggagaa gtaagagcat caagcaagga gtggaacact tttqqttqqq
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```

agtggagagt	cttgatagag	aatactgctg	catcagatgt	ctttttacat	gtgtatttgg	420
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cccagctcct	ggtttgcaga	ctgccacctt	ctcagtgtgt	tttcatgtag	cagagagtga	660
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cagggggcca	atttgggttc	atggaaccct	tggcttccgg	gttggaagga	attttctggc	780
ttaaccttcc	caagaactgg	aaataatagg	gggggccccc	ctgcccggcc	tgattttgga	840
tttttaaggg	aaaacgggtg	ttccccatgt	ggcccagctt	ggctttaacc	teeggeeete	900
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<213> Homo sapiens

<400> 301

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```
<210> 302
     <211> 413
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) ... (413)
     <223> n = a,t,c or g
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acggtcgccg gtgcgaccta ctactaacga ggcagtatgt actgggtcac agtcatcacc
                                                                      120
ctgatctatg gctactacgc atgggtaggc ttctggcctg agagtatccc ttatcaaaac
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cttggtcccc tgggcccctt aactcagtac ttgatggacc accatcacac ccttctgtgc
                                                                      240
aatgggtatt ggcttgcctg gctgattcat gtgggagagt ccttgcatgc catattattg
                                                                      300
ggcgagcgta aaggcatcac aagtggccgg tctcaactac tgtggttact acagactttg
                                                                      360
ttctttggga taacgactct caccatcttt gatgcttaca aacggaagcg ccn
                                                                      413
     <210> 303
     <211> 681
     <212> DNA
     <213> Homo sapiens
     <400> 303
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                                                                      120
tgctgacttc cetttectgg acttgagetg atgaagggga aatggtgttg cagteteete
                                                                      180
tgtcagagcc ctcaggtgca gacggcactt gtctgccccc tcagcctcag ccttggccca
                                                                      240
cetggtecce agtgecetet cetetggetg gggeaggagg acetgeegga catageeaga
                                                                      300
tgtattacgg atgactgcag tcagctcccc caggctcctg cttctcttgc ctcctgcttt
                                                                      360
tttccccaga gctgtctcct tatctccatt cacttgtcta tgggttactc ctggaccctg
                                                                      420
gggttaggag ttggaatcag gctgttaccg acaaaagggg tcaaggtgac tcattttcct
                                                                      480
tatcacgctt aggagttcaa gcgacttgct gatcttccta attcttacaa aacctqccat
                                                                      540
gaacccagct ccctttgtat gactgaccct gccagcctgg gagacataga gtctgattgc
                                                                      600
ccggtctggg ggttataacc ccccggggtt tggacctgga aatccaaagc accctttggg
                                                                      660
gctaagacct gggccaagcc q
                                                                      681
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     <211> 427
     <212> DNA
     <213> Homo sapiens
     <400> 304
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gtcatcatcc gggcaacttg ggcacccacc tcgggctcct cattcatgga gaagatggtg
                                                                      120
ctggtggctc ttcatgctgg ctacatcttt atccagacgg agaagaccat ctacacccct
                                                                      180
gattcactac cgggtgttca ctgtgaacca caagatggac cctgtgacca ggacattcac
                                                                      240
tctggacatc aaggtggtct ttcccgatga ggggtggggg gtggtggtgg atcctggaca
                                                                      300
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```
ctggggttac atggtgtgct gaagtcctgg gggcatgaqc caccaqqqcc ctcccaqaqq
                                                                      360
gcagtcacca gccccacccc ctatccccac agaacccaaa gggaaacacc gtgattagcc
                                                                       420
agagtct
                                                                       427
     <210> 305
     <211> 609
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(609)
     <223> n = a,t,c or g
     <400> 305
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tacccatatg ggtgggaggc tttggatttt tctccagcta tgtcagagcc tgggtctgag
                                                                      120
cacagtggtc agcagcagac ctgttgcctg tctggagtcc gttcctggga tgtgtatgtc
                                                                      180
ggtctgcatg cccttgaatt accgtggaag taacttctct gagacagatg tctggatgga
                                                                      240
tettteeaga geteatettt gaateettgt tattataaaa taagaattaa attgttgaae
                                                                      300
tataacactt agggttaacg ggcacataaa tactttttt aaattttta acatatatat
                                                                      360
atttttttca tcacattttc attgtattag gtatcagaat ttttttttt aattcagtac
                                                                      420
agatttacgg cctggggggg gggctcacgc ttatagtccc aaagttctgg gattacaggc
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gtgcacnctg tgcccggcct aacattaatt cttagttatg tgcacagtct tatgggcaca
                                                                      540
aaagccaaat actctcatgc ctgaagaaag taagcatttt taatgcaaag gtatgagtag
                                                                      600
acaatgatg
                                                                      609
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     <211> 608
     <212> DNA
     <213> Homo sapiens
     <400> 306
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gttcatatct ttcccctttc ttccctgctg ggggatggaa caatgaggct tctaccagat
                                                                      120
atcageteeg aetggetttg ettgaateaa gagtttgeee etgtteaate agecatagee
                                                                      180
atggagtggg ggtcatgtgt gggggatcag gatgacaccc actggatatg tctgaggcag
                                                                      240
accagtgggg tgtaatcact agggacacct acatttgcct gtagtgtaga gagggactga
                                                                      300
tgtcactttg gtgccaggac tgagtggcct tctcaggaac cagagccttt tgccgaaaaa
                                                                      360
aggtttggga tcctgaggcc agaccagtca ggcagtccac cctgaacaga gcccatgcag
                                                                      420
gacagtgggc atgagacccc aaacctctgg ctgagaatat tgccctcact taaagaagga
                                                                      480
gctggaaccc gagtgcagtg cctcacgcct gtaatcccag cactttggga ggctgaggtg
                                                                      540
ggcagaacat ctgaggtcgg gagttcaaga ccagcctggc caacatcatg aggcttcatc
                                                                      600
tctactaa
                                                                      608
     <210> 307
     <211> 781
     <212> DNA
     <213> Homo sapiens
     <220>
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<221> misc feature
      <222> (1)...(781)
      <223> n = a,t,c or g
      <400> 307
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 ctctcgtgga gactagttcc gctgttttgg tgcctgcaga gcctcactgg ctttctaggg
                                                                       180
 ccctgcttgc cacgcaccac acgggcattc ctctctctgc agtcctggga cctccctggg
                                                                       240
, actogaccag gaagccaggo acagggetto actgottgca atgotgcaaa cacacctggo
                                                                       300
 ttggcggcct tgccaggctc aggcgctttc tctgtgatac cagtgtcctt gttattgcct
                                                                       360
 gtaccagagg ggttgggtag aacttacctt tattcgtgat gtttcagatc acatttttta
                                                                       420
 tccatggcta tgagtccttt ccattcttcg aggatcctgg attctgaaat tcaaaagcca
                                                                       480
 gggagaggcc gggcgcggtg gcttatgctt gtaatcgtag cactttggga ggctgaggtg
                                                                       540
 ggcggatcac ttgagcccag gagttcaaca ccagcctgag caatatqqcq aaaccctqtc
                                                                       600
 tctaccaaaa atacaaaaat tagccagcca tggcggnggg caactgtaat cccagctact
                                                                       660
 cgggaggctg aggcaaaaag gtttgcttgg acccaqqaqq caaaqttqqc qtcaqccaq
                                                                       720
 aacatggcac tgtactccag cctgggcaac anagtgagac cctttttttc caaaaaaaa
                                                                       780
                                                                       781
      <210> 308
      <211> 1391
      <212> DNA
      <213> Homo sapiens
      <220>
      <221> misc feature
      <222> (1)...(1391)
      <223> n = a,t,c or q
      <400> 308
 tttacaacca actitittit tattittit tttaaattit tcattitati caaagtiggt
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 acagaattgc taacatttcc ataaaataat tactatactt cagttacagg acaaaatacc
                                                                       120
 acagaaagga atgtactttg caagaaatgg tagttcatcc taagtttcca aatacttttg
                                                                       180
 gaaggctaat gcagcagctg ggcaaaataa cacacagtac acaaagaaca gtgtatttca
                                                                       240
 cagagtcagt aatgaaaaac tgacagctct ttaggcagga tatgcttttt ttcattttt
                                                                       300
 taaacaataa ccactttcaa aaacacatgg aaccaagatc atacatggtt ttacaatttt
                                                                       360
 aaaaaatcag attgtacaca ataggttaga atagacaagt tagaattgtc atgattttaa
                                                                       420
 caatcttaaa tctacaattt caactgtact cctttcaata tagaaataac ctgctttata
                                                                       480
 ccaaattcta ctttctgctt gcaactaaaa cactgtacaa tgagatggat acaattagtc
                                                                       540
 aaaccttaaa attaaaaaag ctgtagacaa cagaaggtaa actggaaatc catttacaat
                                                                       600
 tcaaaaaact cactaataac aaaattaatg ttcatcaact tcatttataa tcacatttgg
                                                                       660
 cctacaatgc ctaactaaaa tgacacatgt acacaatata cacccccagt gtactaactg
                                                                       720
 gtctcttaca aaaaatctga acaaagcatc ataagcagga cactgggaag aacatgtttc
                                                                       780
 aatgtagaca tettttaaaa atgeattaat aettaeatat caaaattaet agataaaage
                                                                       840
 agcagcactc tgctgacatt tggcttaaaa ataaatgaat gaatgaagca atttcacagg
                                                                       900
 atattattag aaaaagaatt ggttttcttc ttgaagaaga ctactaactt ttgcacaqca
                                                                       960
 actatttttg atatccatct tatcaaaaag aaaaaagaaa gcactgagaa gtataacaca
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 gttcatacat gattgccaac atgggtctgg acaaaagaaa atgggatgtc caagcaaaqa
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 acgggtaaat ccctgctcta tttctgaact ctgctggcaa tctataaact gaagcagtaa
                                                                      1140
 cagtggggga aagcaaggga acaaattcca taccatcatc tgacactaat ggagtatggc
                                                                      1200
 attattaaaa aaaataaagc ttttgcattt taataacccc acagaaaagt ctatgagcaa
                                                                      1260
 aagacttgat ctgtttgcca ctcaaaagtt agagatctca cagtgaaatt agaaaactct
                                                                      1320
 aattatacat atttcggacg cgtgggtcgn ccctgcagat ggngatcatn ccgacgggat
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 cagtgggggc c
                                                                      1391
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<210> 309
     <211> 874
     <212> DNA
     <213> Homo sapiens
     <400> 309
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ctgactttat accttcattt cagcgtggta aaaatcgatt aacacttcta atgagtcaag
                                                                      120
tcctagggtt ttttggtttt gttttgttgc caacgaggaa cacagctctg ggggaatggt
                                                                      180
gtcatccacc tcgctttaaa aataagcaca tgatggctgg gcaccgtggc tcacgcctgt
                                                                      240
aatcccagca ctttgggagg ctgaggcggg tggatcacct gaggtcggga gtttgagacc
                                                                      300
agcctggcca acatggtgaa accccatcgc tactaaaaat ataaaaaatt agctgggcat
                                                                      360
ggtggcgcac gcctgtagtt ccagctactc aggaggctga ggcaggagaa tcgcttgaac
                                                                      420
ccgggaggtg gaggttgcag tgagctgaga tcgcaccatt gcactccagc ctgggcaaca
                                                                      480
agagcgaaac tetgteteaa aaaaaaaaaa aeeeeaeeee caaacagaaa aataataaag
                                                                      540
taacttcaga attttaatgc tagaaattaa aggtagcatc cacacataat tccacctgca
                                                                      600
aaatetttag tgagaagatg acaatacgat ettaetecaa caqttecaat eetaaaagae
                                                                      660
atccaaatta tgataaattt tagtcttatg aatgcgagga aagggtgaaa agaggtgctg
                                                                      720
gaaatacagc atgcagacca aacaaaaatc tccacaqtca ctqaactcat attctaqtat
                                                                      780
agggagcccg aaaacattta caagtgaatc tacatcactt tgatagagta agaaggcaag
                                                                      840
tgggaattcc gccacacgaa ctagggatct cgat
                                                                      874
     <210> 310
     <211> 802
     <212> DNA
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     <400> 310
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                                                                       60
cacactetta atagcaaage aattttggat attcaccgtg qacctacatt tgtcagatta
                                                                      120
tgttttggag ttatctaggt acctaataaa tqcctqtttt tacaqcccat qttcacaqcc
                                                                      180
cattgagaaa tagacaaagt gggtaaggca gatgaatgaa aacatgtcag ttttattact
                                                                      240
gataatgtac tgcaattgga gaatgtggtc agatattcca aacttcctat gactgcacac
                                                                      300
tgaagagtct tctctttgga ggggagaaaa ataatgctcg tggctgtttt taaaattatg
                                                                      360
tttattatat atttattaaa agaaagataa tatttagaaa aaaatctcat tagtcaagta
                                                                      420
aaattttaga tactctatct tgaaaaacct tctgaaaaca gtataaaaaa tatttgagat
                                                                      480
atgtcagtat aacatagagc aatattcgat tctccctcct tggggcagca aatatttct
                                                                      540
gaaaatcaaa agtacagaat cttttaggca ggaaatacat tttggccaat tataatttta
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gaagtcaaaa ttgttaaggt ttttggacca agcacaatgg ctcacgcctg gaatcccaac
                                                                      660
actttgggag gettgaggea ggeaetteae ttaaggteaa gagtteagaa eeageetggg
                                                                      720
caacatggtt taaccccccc ctcccttaag cattacctaa tttattgggg catgggggaa
                                                                      780
cactacgcct gaaaccccag cg
                                                                      802
     <210> 311
     <211> 352
     <212> DNA
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     <400> 311
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gegaacagae etgettgete agttgetgtt tttaggaaga ggtgateece gtaggagate
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tgaccaatgg ccggacacta taacttgaag ctgccaatta ttqcaqcaca tgggactggt
                                                                      1.20
aacaggagca ccatttcctt gagctcctcc acgccaaggc ctgtgagcac catggggagc
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aacaccttta ccaccttcaa tacaagcagt gctggcattg ctccaagctc taacttacta
                                                                      240
agccaagtgc ccactgagag tgtatggatg ccacccctgg ggaatcctat tggtgccaac
                                                                      300
attgetttee etteaaagee caaagaggee aateggaaaa aactggeaga ta
                                                                      352
     <210> 312
     <211> 1267
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(1267)
     <223> n = a,t,c or g
     <400> 312
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gcaaaagtgc aaatttgatc atgctgttct tctgctccag atttttcagt ggcttctcaa
                                                                      180
ctcattcaga gtaaggccaa aatccttacg aagtcctata atcatttgaa tgatctgttt
                                                                      240
ttgtctgcct gtctgtccta aaacacacct ggctcatccc atgctagcaa cattggcctt
                                                                      300
tgtgtcactt cttgaatatg ccaagcattg cctcagggac ttcatacttg tgtcctttct
                                                                      360
tettggaatg etetttetea gatateaaca etaaacaeta eeacteetea aatateaeta
                                                                      420
aatcactaaa tcaatcctgc cttatttaaa gagaaatctc acttctctct gcagttttaa
                                                                      480
atttttttta gattttattt taggttcaga ggtatatgtg caggtttgtt atataagtaa
                                                                      540
attgcatggc atgggaattt gctgcataaa atatttcatc actggggtga taagcagaat
                                                                      600
acctgatagg gaactttttg atcctcaccc ccctcctgcc ctccgtcttc aagtgggccc
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tggtgcctgt acctcccttc tttgtgccca tatggattta aaggtcacct cccacttgga
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agtgagaaca tgtgggcctt gccttggtgg tccctggccg agccttcgcg accacgggaa
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ttaaaacagt gtcttttctc tcaccgtgag aagcctgcaa actgccggtc cgcgaggggg
                                                                      840
gcgccctgtc gcatgccgac atttggggaa ccgcgcatca acaccttacg ccgaatctcc
                                                                      900
gcacactacg cgacagtgag acategtega etteceeega taegeggate tegeegagte
                                                                      960
gcgtcgcact ccgcggctca ccgccacgtt ggccaaccgg tggcgacctc cgctatggtg
                                                                     1020
acgacctcgg cattttctgc gttcctcgct atcccaccgc cctgtgggaa aactccggtc
                                                                     1080
gtccggcgnc cggcgcggtc tcacctataa cgtcccgcat acgccggaga gacagaccta
                                                                     1140
taacctcgca tattcgcgcc atccgcgcaa ttcgcacgca aaccgatcct aaccacccgc
                                                                     1200
gccatcgcgc gcgattccaa ctgcgctcgt ggccctaggg cgcgggaaac tccgcggctt
                                                                     1260
cgcgtct
                                                                     1267
     <210> 313
     <211> 1927
     <212> DNA
     <213> Homo sapiens
     <400> 313
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aatgtgttaa ccaaaagcat aatatattcc cagtaaacaa ggacttccaa cttatcctat
                                                                      120
aactaaaaag tcaactaaac agttggtttt agctagagac aaacatcaqt cactqccacc
                                                                      180
aaattccatt atataaattt attttgcttc acatttaagg agaaacccag cagaggggtc
                                                                      240
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<213> Homo sapiens

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                                                                      360
ggcctgcagg ctaggatggc ccggccctca gccttcccca tcgqqgtctg tctqactctq
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eccatggeet ggateteece gggtttaget gtgeecaget gteeceagta cataetteaa
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                                                                      780
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gggattgtga agcagcagat agacagccat atcacagatc cagatcaaca gaacaacggc
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                                                                      366
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     <211> 839
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tgtgtacaga tcatagtctg aagtggaata agcagaatgt tgtcctcagt gtgagatgtt
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aacaggtact caagacctgt ctgggctttg gcctttgggc acattccccc tcatcacctt
                                                                      360
cetteceact tggetgaget atggatgaga aaacetaggt caatagttea ceaacteace
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                                                                      480
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                                                                      540
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                                                                      600
aatttgagac caacctgctt gggccaccta aacccatttc atcaatcaat cataatcgag
                                                                      660
ggaggggggg gattggagcc ctcattatta ggagctgagg ggggggccac tggaccccgg
                                                                      720
ggtttgggtt gccgggcccc tattggcccg gaccctggga aaaaacgaaa accagcctcc
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                                                                      240
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acceptatec acaggaacag taactatagt ttgtcctaat ataacgaagt ctactttata
                                                                      420
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                                                                      540
aactaataat aaaaaaaatt agctgggcat ggtggcacac gtcctgtagt cccacctacc
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ategeaceae tecaete
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aaacaagccg ggtggctgag ccaggctgtg cacggagcgc ctqacgggcc caacaggccc
                                                                    180
atgetgeate cagagacete ecetggeegg gggeatetee tggetgtget eetggeeete
                                                                    240
ettggcaccg cetgggcaga ggtgtggcca ceccagetgc aggagcagge tecgatggcc
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ggageeetga acaggaagga gagtttettg etectetece tgeacaaceg eetgegeage
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tgggtccagc cccctgcggc tgacatgcgg aggctggact ggagtgacag cctggcccag
                                                                    420
ctggctcaag ccagggcagc cctctgtgga atcccaaccc cgagcctggc gtccggcctg
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     <211> 992
     <212> DNA
     <213> Homo sapiens
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cttgttttag atattttact gcttcagaga qqqtcatqtt cacaccattc tccccttttq
                                                                    420
taatttttca caceteeetg geteecettt tataatttag aaagaggttt acaagtetqt
                                                                    480
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gttgctgaat gactggaaac ttgtgtttct tttccattaa gggctatttg ctgacttctg
                                                                    600
aaatattgat gatttatttg actttagaat tttgcatact gaggggaaag catcttaatg
                                                                    660
tatcatttaa agcaggagat actttcatac tatacctggg ttctcttggc tttgaagagg
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aagctgagge egggtgeate atttgggget taggagtttg ggaccacccc tgggccacca
                                                                    840
900
ctatacatec agttteteet caggegggee cattatatta aaccetagee ggeegeteee
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tegeceege geaacaatat atetateege ee
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accagagaga tetteeetga eeatteaata teaaatatta eteettetgt tacagtaggt
                                                                    300
agetagteag geatgageag ggeagaagag ggeteeeete eeteaacaca caccaggaat
                                                                    360
gacaggcaaa catcaggtga tggtcaggca gctgctaact gtttctctaa aatattaatt
```

420

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ggttgcagcc tgcaccaggg aaaggcagtc tccatatata cagaagcacc tgaagctggt
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gaacacctca agtgaacacg cgtacaactc cagtaaacac gttgcacatg gtccctttcc
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caagtgctgg gaggctactg tgtgtgcaga cagcctgccc caagggaaga atcatgggag
                                                                      720
atgggacacc aagateetgg aagtatgeea acatataaaa ceecaagttg aaaggteaaa
                                                                      780
cogtgoattt gtottttcaa gttgcccact ttgccctctt ccaagtgtac cttccttccc
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gagetatteg atagtgaaga eeceeggeag egagagtaee ttaagaacat eetgeaeegg
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ctttatggca ggatgctggg actccggccc tacattcaca aacagagcaa gcacattttc
                                                                      180
ctccggatga tctatgaatt ctagcacttc aatggggggg ctgaactgct ggagaaccta
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ggaagcatca tcaatggctt tgcgctgccc ctgaagacgg agcacaagca gttcctgggt
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cgcgtgctga tccccctgca ctctgtcaag gcgctgtctg tcttccatgc ccagctggca
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tactgtgtgg tgcaattcct ggagaaggat gccactctga cagagcacgt gatccggggg
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                                                                      423
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332

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420

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		aggttggaaa				3900
		gaggccaggc				3960
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120

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360

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638

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2940

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435

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                                                                     1380
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360

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420
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gegagtagag getggtgegg aacttgeege eeccageage geeggeggge taageecagg
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<213> Homo sapiens

<400> 400

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cccttcgctc agtcccggcg cctttaaagt cgccttccaa aaaattcact ccccagccac
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     <212> DNA
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                                                                    420
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cccggtggct catgcctgta atcccaacac tttgaagagg ccacggtggg tggatcacga
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tgcagtccta tttctcctcg cgcctcggcc agccactgag actccttgca tcagataacg
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     <211> 554
     <212> DNA
     <213> Homo sapiens
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cagagaacaa agtcctgcgg gaagagaatg acttggaagc cggcaatctt catcctcagc
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aggatcaaag ctgtctcaag gagtgccctt gcatgaaagg aggcacagat atgcagacca
                                                                    240
agaaagaggc aagtgctgag acagaatata tgaagcaaca atatgaagaa gaccttcgta
                                                                    300
aaatcaaaca tcagacagaa gaggagaaga aacatctcaa agaccagcta gtgaagcgac
                                                                    360
420
ctgaaagaaa gaaactgcag agggaagtag aagcacagtt ggaggaagtg aggaagaaat
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cattaatcgg tatttgaacg tgattttaag taattatgtc taaatacagt ttgttcagtt
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atttgagget acattttata attaateeea tetaaattta ttttgteaet gtttgagaet
                                                                      240
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                                                                      300
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                                                                      360
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                                                                      420
agttattctt ttgttaaaag agtataatac tgtttttgag agaatatgat atgattccat
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gcaattcaca totgtgttgc agttagattt aattatttgg actgggaagc cccatattaa
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ccagtgcaca tatgtagtcc cagctgctcg ggaggctggg gttggaggat cgcttgggtc
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caagaggtgg aggttgcagg gagccatgat cacaccactg tactccagcc tgagtgacag
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ctgagtaggt tctgcgaagc gatagcaacc gccaccgcgg cggagcaccg ccctcccta
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gctgtgggaa gaggccqaqc qtqqctttat cttqcactca tqcaaaaqaa actqqcaqat
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tatctgaaag tgcttataga caataaacat ctcttaagcg agttctatga gcctgaggct
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egeggetggt ceteagtgea cacetgagta geacgacett teegecetgg acgeacgetg
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acaaaacatc agtgaagagc tccagagaaa tatttctcta caactgatga gtaacatgaa
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1281

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<213> Homo sapiens

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                                                                      660
getttttet ttttaggggg ggaattttte tttggeeceg geegettttt aaacggggga
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aa 1322

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acceptinged to to to to to to to the total accepting the control of the total accepting the total accepting the total accepting the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total accepting to the total acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance acceptance accept
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aatagcagag aaatgtgatc gctgagatgt aaaaagtttt taatgctagt ttccaccatc
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540

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328

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332

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		gagcaccctt				660
		ttaaaagtag				720
		aagtacctcc				780
		ggattaacag				840
		cggaaggagg				900
		aacaatgggg				960
		atcctacgag				1020
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		gacaggctgg				1440
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		cctccggcag				1560
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		ccagatgagg				1680
		atttctgagg				1740
		atttcctgta				1800
		ttcttccacc				1860
		ccatctctca				1920
		aggggcagct				1920
		acccctaaca				
		tggagaaatc				2040 2100
		gaaaaatact				
		atctggtacc				2160
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		aatttccctt				2340
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		tagacatatc				2880
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		catttacccc				3000
		tctgtcccac				3060
		atgttggagg				3120
		ggttgctgcc				3180
		ggtagatcag				3240
		tgggcaggtg				3300
		ttgattgatt				3360
		ccagaggaac				3420
		aggtcatgtc				3480
		aagcccaagt				3540
		ccacgctgct				3600
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gcaatccttt atggtagata atgttgtccc tacattgtat acaagaaaca aaggtgtagg
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                                                                         780
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  geoceaecge ggecateaeg eccategege acagegteet ceageegeeg eccetettge
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gctttgctcc accgcccgct accactgcaa gaacggcctc tgtattgaca agagcttcat
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ctgcgatgga cagaataact gtcaagacaa cagtgatgag gaaagctgtg aaagttctca
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1080
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                                                                     1902
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<211> 2122
<212> DNA
<213> Homo sapiens

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tctatggggg ggggggggg cg
                                                                     2122
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ctgagtgcca gtgcccaccc cacatgacag ggtcccggtg tgaggagcac gtcttcagcc
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gatgtgagga gtgatgacga ggattatgaa gaqqaaqaqg aaqaqqaaqa aqaaqaqqct
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accaaaggca aagagatqqa ttqtttaaaq aacqqcctcq qqqctqaqaq qcacctcatt
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ctctttctca ccaccgcctt ccctcccacc taaqatqtqt ttaccaaaat qttqttaact
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tgtgttaaaa tgttaaatat aagcatgccc atggattttt actgcagtta ggactcagac
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aagteeteag tgttteeetg geacaagaac eteetggeag tagatgtatt tegtteacet
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				ctcccctgag		2760
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				ccctgcgccc		3000
				ccccgagaa		3060
				gcttggagct		3120
				gggacgccat		3180
				ctctggtgtt		3240
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				gcagttcccc		3360
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4179

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<211> 2220
<212> DNA
<213> Homo sapiens
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<211> 2095
<212> DNA
<213> Homo sapiens
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PCT/US01/02687 WO 01/54477

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360

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360

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<213> Homo sapiens

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<213> Homo sapiens

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<213> Homo sapiens

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gtcagcatat agggacagaa tataactaca cattaataat ttctcaagta tttattttag
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     <223> n = a,t,c or g
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397

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<213> Homo sapiens

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aaaa
                                                                     1024
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380

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:· ·

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240

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gacaaattat aataaattat taaaagaget ataatggata taaagtgtgt gttctgacag

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1380

1421

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actggggggt tetgggtgtg attgtgegte tettgttttg atcagaacee acttagggee
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eggggtetgt tgteccetee agatggagga cagggatett ttecacetea cetgtgtece
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cagtgcccag tacaggccca ggcacaaata ggcccttact tcagagaact gggtgaacca
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                                                                      720
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                                                                      780
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gttgcttgag aggctgaggc tggaggatca cttcagcccc ggagctcaag gttacagtga
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teettettgt aeteeteeaa teteattagg ggeeggaagt agatgggata gaaggeggeg
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gtagccagaa gggttttcct ctttgcctca gttagtaccc agggcaaaag cttaatgtat
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tctactcaqa aagtaqttaa ataagactgt ttctctaata tatattttag ttgtaggaat
                                                                      420
                                                                      480
taggaagtag catcatagat geteetacae taagetggee etgetteeta tgttaaatat
gacacatetg aggeeetggg agaggaagtg atttgeeeag teteacacaa tgagttagag
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ccagagtgaa gtcaaaaccc agtctctgga tgtacaagca aggtcttttt ctagtcccaa
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gaaggtctga tctttgagtc agagtcacag aagaattgag aatagctgtt gggccttggg
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gg
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tactttagcc acacgeggaa gaacageeca ctacattact atcagegtet egagategte
                                                                    240
gaageegeaa ttaggaettt gtttteegte aetgggatee tggeagagea gtttgtteeg
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gatgggcccc acctgcacct ctaccatgag aaccactgga taaagttaat gaattggcag
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cacagcacca tgtacctatt ctttgcagtc tcaggaattg ttgacatgct cacctatctg
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gtcagccacg ttcccttggg ggtggacaga ctggttatgg gctgtggcaa gtattcatgg
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aaggttteet ettetaetae caegteeaea aeeggeetee getggaeeag caeateeaet
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cactcctgct gtatgctctg ttcggagggt gtgttagtat ctccctaaga ggtgatcttc
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cgggaccaca ttgtgctgga acttttccga accagtctca tcattcttca gggaacctgg
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     <210> 649
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gagactttgt ctcgaaaaca aaagatactg gggccatagg aggaatgtga taaaccagat
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acggggtttc accatgttga ccaggetggt ctcgaactcc cgacctcagg tgatccacct
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gcctcggcct cccaaagtgc tgggattaca gatgtgagcc accatgcctg gccctgtttt
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                                                                      600
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                                                                     1080
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ggga
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     <212> DNA
     <213> Homo sapiens
     <400> 651
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ggcacaaatt tattttccta ttctgcagct cgccttttcc cattctgtat tttcctagtc
                                                                      420
ctagettate titteteatt etggattiet tettititga eatggageet eegettitge
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gtecaagetg ggeggegtgg eeeggacetg ceteactgea atgteegeet geeaggtgta
                                                                      540
ategetttet cetegeteea ceetgeggt agttegagge teactgettt aacetetege
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<210> 652

<211> 743

<212> DNA

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atteatttea aaggaettte taattteeet tgteatttet tetttgatee gtgagteett
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cagaagggtg tagtttaatt tcaaaatatt tgggqatttt tcaqacactq attttctgtt
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aacttttggt ctatctaacg gaatgctgta tggcacttga aqaaagggtg cattctgttc
                                                                      420
ttatagggtg gagtgtttca tttaaaaqaa tacaaaqqca attaaaccaa qtqqqcttqa
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tagagttett caagatggte etetgeagea acacagatgg aactgaagge cattateeta
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taaacaatgg gtacacatgg acatagggag taaaataata gacactggaa actccaaaag
                                                                      660
gcaggaggat gggaggag taagccatga aaaatcacag attgagtaca atgtacacta
                                                                      720
aaagcccaga gttcaccact atg
                                                                      743
     <210> 653
     <211> 1524
     <212> DNA
     <213> Homo sapiens
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gaagacaatg aggcaatete agcaetttgg gaggeegagg etetetgttt eetegagtea
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ccggcccgga ccacctcagc ctgctgtgcc tcttcattgc cagcaaccgc gtcttcactg
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agggeacetg gaagetgetg acet
                                                                     1524
     <210> 654
     <211> 711
     <212> DNA
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tccctggggg ccataattgc aaagtttatt aatatattat cctatatgta ttaatcctgt
                                                                300
aggtcctaag gaaataattc aaatttgggg aagggaacaa agctctatgc ataagatttt
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aagacggtgc atccatagaa ttggtggatg aagagccatt gaaaatgatg tttgggggcc
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aagcatggtg gctcatgcct gtaattccag tgactcagga agctgaggtg ggaggattgc
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tttcaaaatt agctaggtgg tgcgggccta tgcctgtagt cccatctact tgggaggctg
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```

<210> 655 <211> 1524 <212> DNA

<213> Homo sapiens

<400> 655

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<210> 656

<211> 993

<212> DNA

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<211> 394

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gctgtagtgc agtggcacaa tcaccactca tcgtagcctc aacttcccag gctcaggtga
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                                                                     1121
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cacagetaat eggeattate actateteta ettetateat aacaaeggtt acegeegtgt
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360
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caqaaatgtq aacaacaaaq atttttttqa tccaqttqaa aqtqatqaag acatagcaag
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geggtgacac aattataget eagtgeaace teaaatteet gggeteaage tateeteeca
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                                                                     480
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cqcctqtgqa ccttttcaaa tatggctggt gtggagctgt gcccagggcc ccagccagcg
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ggtcctgctg cccctgttgg gaggacgccg cctgtcctct ctgctttcac aacaacctct
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tattttgaac atcagcagct gaggcaactg aacatgtttc tgtgctgtct tgcacccact
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tetetttgga agetteetat gtattaetge acacetttte catgeeteet etgteeteeg
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cttcaacctt ccagagatgc tccagggtat cagtgggtcc catggaagac tgtctgaacc
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aagacaagat aagatggaaa gcctcccgaa agacatgggt aggttcttag atgaacaatg
                                                                      360
ggtttatttt attattttat tattattatt tttttttcga gacagtctcg ctctgtcgcc
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tgataaaact ttagactcta aaaagattgc acacttcaga gctgagaaag agactttcag
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cgaaaaagat acatattgct atttaaaaat ggaactctga aaattaagca tctgaagacc
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cagtggcgtg ttccatgctg ctgtaggcca ggaacatggt gcagccgaag tggacggcca
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tccagtgatg acttggcccc agtggacagc tgcccagtga tgggacatct ggagtagatg
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actagagete egetecattg ggttggagee attecagggt gggaatggee accaggagae
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gatgcctacc cttctcttct tgcaccaagt cagcacccat actcaggcga ggccctgtgt
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gaggatatta accetttace ceattgttgg gaacgggatt tgggggaata aaaacttttt
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gegeetteaa gaaacteaag aatttgaaca caetgtaeet gtataagaat gaaateeatg
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tgtgattett eccetettgg ggetgetget etecetecce geeggggegg atgtgaagge
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cccctaccag gagatcgcag gggaacactt aagaatctgt cctcaggaat atacatgctg
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                                                                      360
atcaatgate tecageetga ggatgagtet gaatattaet geettgetat gacageagee
                                                                      420
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447

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                                                                      180
                                                                      240
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120

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531

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420

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                                                                      840
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                                                                      120
tettgegetg tgteeceate tetaateact tataetteta ttggtacaga caaatettgg
                                                                      180
ggcagaaagt cgagtttctg gtttcctttt ataataatga aatctcagag aagtctgaaa
                                                                      240
tattcgatga tcaattctca gttgaaaggc ctgatggatc aaatttcact ctgaagatcc
                                                                      300
ggtccacaaa gctggaggac tcagccatgt acttctgtgc cagcagtgaa agggggtctg
                                                                      360
gggccaacgt cctgactttc ggggccggca gcaggctgac cgtgctggag gacctgaaaa
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acgtgttccc accc
                                                                      434
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                                                                      120
tgtagccaca gctgaggccc tggaccagct ctctccacac cgcatgctcc gagttgggac
                                                                      180
tctaaggagt ctaggaattt tcattcaaac ttggccttac aggtcactca tcagaaaaat
                                                                      240
acttttttca aggtcaacca atagaacata ctttattcaa cagtttgtta gtttgctttt
                                                                      300
taaatattta gccacatggt atgtaggett ccatgtacac tettgecetg gcccetqaaa
                                                                      360
cataagcagg gggctcttct gtacatttgc ccagcttccc tgccagcctt taaccccagg
                                                                      420
aaceteteag tetaceteet ettttetgee tetgaateee tacetttaaa gteagaacag
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gccaggcccg gtggctcacg cctgtaatcc cagcactttg ggaggctgag gtgggtqgat
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cacttgacat cagtagttca agaccagcct ggccaacatg gtgaaacccc atccttacta
                                                                      600
aaaatacaaa aattagecag gtgtggtggc gggcacctgt aatcccagct actcaqqaqq
                                                                      660
ctgaggcagg agaatcactt gaacccagga ggcagagttt gcaqtcaqcc aaqatcacqc
                                                                      720
cactgtactc cagcctggat gacacagcga gactccgtct caaaataaat acaaaaaaaa
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aaaagg
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ccaccatcga ctgcaggtcc agccagagtg tcctctacca cgccaacaat aaaaactact
                                                                      180
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taacttggta ccagcagaga ccacgacagt ctcctaaagt gctcattttc tgggcatcta
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cccqqqaaac cggtgtgcct gaccqattca ctggcagcgg gtctgggaca gattattcgc
                                                                      300
tcaccataag cagcctgcag gctgaagatg tggccactta ttactgtcaa caatattatg
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attctccgat caccttccg
                                                                      379
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                                                                      120
ccaacettet caaceteetg etgetgtett tgtttgeegg attaaateet tecaagaete
                                                                      180
acattaatcc taaagaaggg tggcaggtgt acagctcagc tcaggatcct gatgggcggg
                                                                      240
gcatttgcac agttgttgct ccagaacaaa acctgtgttc ccgggatgcc aaaagcaggc
                                                                      300
aacttegeea actaetggaa aaggtteaga acatgteeca gtetattgaa gtettaaaet
                                                                      360
tgagaactca gagagatttc caatatgttt taaaaatgga aacccaaatg aaagggctga
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aggcaaaatt tcggcagatt
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     <212> DNA
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     <400> 747
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tggaaagaaa aataaccata tatacaaaat catgcataag aaaaaaataa tataaggatg
                                                                      180
tacataccaa atattaataa taatggctat ctctggatag tggaatcaga gggattatgt
                                                                      240
aattttcctg ataaattttc ctgtcctcca aacagcatcc gcttcatact attatttctt
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ggttgtaatt agtttgatat aattetette agaaaggete tgttteaeta tatatacete
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aaagcatact tttgatgcag cttctgcaat tcccatctaa aaagtagata acacttgctc
                                                                      420
ttatattctg gcatatgaag actatttgta attaacacac tataaaatat gtcaaagcag
                                                                      480
gccaggcatg gtggctcaca cctgtaattc caaaaccttg gcaggaagat cgattgaggc
                                                                      540
caggagetea agaegageet gggeaacata gaaagaeeet atetttacaa aaaaaaettt
                                                                      600
aaaaattagc caggtqtaat agcacatgcc tgtctgtaat cccagctact tggcaggctg
                                                                      660
gaagqtcaaq qctqcaqtqa gccatgatca tgccactgca ctccagccta ggtgacagag
                                                                      720
caagaactca tctctaaaaa aaaattttta aataaagcaa aatatgccac agcatagatc
                                                                      780
tgattgtaga aaattattat atggagaact gaaaaatctc ctaatcaaga caaaaatttt
                                                                      840
aaataqaqqa aaaaaatact atctatcatt agttcaagtt tccattaaga gtagagtgtg
                                                                      900
aagtagetee aagtteagag etggagaatt ttgeatetet ee
                                                                      942
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     <211> 1050
     <212> DNA
     <213> Homo sapiens
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     <221> misc feature
     <222> (1)...(1050)
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<223> n = a,t,c or q

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aggacaccag attgtataca ctgtgatcaa aaccatgtga aaaacacatg catgaagagg
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actgggaaga aatacacaag aagtggttgc attagggtga gaaggagtat tcatgttttt
                                                                      240
ctcatccgtc tttttcaaac cttttgtaat gggtggtttt attaatttta taatggaaaa
                                                                      300
tgttaattta aaagcaagtt atttacagtt tagtaagctc atggcaggga aaggctgggc
                                                                      360
tctgtttatt gctcttactt tttcccaacg cctactccca tgcctggcaa ttatagagat
                                                                      420
aataaatgtg ggtgtggaat gagtgcccac tgggaaacct ctcagaggac tttgacccag
                                                                      480
gaacatattt gcacagggtt tccctcagct ggagaaggtt tctctgggag agcaccagcc
                                                                      540
aggtgtgtgt catgggatat atttacaggg tggtgagctc tcctggtcca acctaaaagg
                                                                      600
teccageaag gtgtagggge eettetggee atttgaeate accagggeag ttagtgetga
                                                                      660
tacaaaccac agagaatgaa caaactccaa ctcaaacggg aatggatttt atgtcattct
                                                                      720
gggactttca aacttgataa tagaccaagc atggtggctc acacatgtaa tcctagcact
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ttgggaagcc aaggtgggag gatcgcttgc ggccaggaga ttgagaccag cctgggaaag
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gtagcaagac ccagtctcta caaaaaaatt ttttgttctg ttttgttttt gagacagagt
                                                                      900
ctcaactctg tcgtctaggc tggagtgcag tggtttgatc ttgggtnatt agtttctttt
                                                                      960
tttgtgggtg ttgtgtttaa gtttttgttt tgggttaaat taatctggtc ttgggaatcc
                                                                     1020
ttctttttat cgttggtgga gatttaaccg
                                                                     1050
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     <211> 390
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
     <222> (1)...(390)
     <223> n = a,t,c or g
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                                                                      120
acagacagee aettteacet getetggaga tgaettgggg aacaagtata tttgttggta
                                                                      180
tetgeagaag ceaggeeage ecceeqtqqt acteatqtat caaqataaca aqeqqeecte
                                                                      240
agggatecet gagegattet etggetecaa ttetgggage acagecacee tgaecateag
                                                                      300
cgggacccag gctacggatg aggctctata tttctgtcag gcgtgggaca cgaatggagc
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tgtgttcgga ggaggcaccc agttgaccgn
                                                                      390
     <210> 750
    <211> 441
     <212> DNA
     <213> Homo sapiens
     <400> 750
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cttgaaggga atccaactta gctttaatta acattcttaa ccttcttacc tctctqqatc
                                                                      120
tcagttgtct catctgtaaa aaggagataa aaattattta cctgcctgaa catgaggtgg
                                                                      180
aggaccatcc tgctacagta ttgctttctc ttgattacat gtttacttac tgctcttgaa
                                                                      240
gctgtgccta ttgacataga caagacaaaa gtacaaaata ttcaccctgt ggaaagtgcq
                                                                      3.00
```

```
aagatagaac caccagatac tggactttat tatgatgaaa tcgttttaga agagcttggt
                                                                      360
ggtccatgcc tatatcttga agggaatcca acttagcttt aattaacatt cttaaccttc
                                                                      420
cgcacgcgtg ggtcgacccg g
                                                                      441
     <210> 751
     <211> 449
     <212> DNA
     <213> Homo sapiens
     <400> 751
gtggggaatt ccccagcaat cagactcaac agacggagca actgccatcc gaggctcctg
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aaccagggcc attcaccagg agcatgcggc tccctgatgt ccagctctgg ctggtgctgc
                                                                      120
tgtgggcact ggtgcgagca caggggacag ggtctgtgtg tccctcctgt gggggctcca
                                                                      180
aactggcacc ccaagcagaa cgagctctgg tgctggagct agccaagcag caaatcctgg
                                                                      240
atgggttgca cctgaccagt cgtcccagaa taactcatcc tccaccccag gcagcgctga
                                                                      300
ccagagccct ccggagacta cagccaggga gtgtggctcc agggaatggg gaggagqtca
                                                                      360
teagetttge tactgteaca gactecactt cageetacag etcectgete actttteace
                                                                      420
tgtccactcc tcggtcccac cacctgtac
                                                                      449
     <210> 752
     <211> 524
     <212> DNA
     <213> Homo sapiens
     <400> 752
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ctaggggttg gcaccggccc cgagaggagg atgcgggtcc ggatagggct gacgctgctg
                                                                      120
ctgtgtgcgg tgctgctgag cttggcctcg gcgtcctcgg atgaagaagg cagccaggat
                                                                      180
gaatccttag attccaagac tactttgaca tcagatgagt cagtaaagga ccatactact
                                                                      240
gcaggcagag tagttgctgg tcaaatattt cttgattcag aagaatctga attagaatcc
                                                                      300
tetatteaag aagaggaaga cageeteaag ageeaagagg gggaaagtgt cacagaagat
                                                                      360
atcagctttc tagagtctcc aaatccagaa aacaaggact atgaagagcc aaagaaagta
                                                                      420
cggaaaccag gtagtctgga cattttcctt gctttttgat ttatttaggg gacaactgaa
                                                                      480
aattttaagc taatgaataa agaggctgaa gaagaaaaaa aaaa
                                                                      524
     <210> 753
     <211> 474
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
     <222> (1)...(474)
     <223> n = a,t,c or g
     <400> 753
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ategeceteg geagectegg cetecacace tggeaggece aggetgttee caccateetg
                                                                     120
cccctgggcc tggctccaga cacctttgac gatacctatg tgggttgtgc agaggaqatq
                                                                     180
gaggagaagg cageceect getaaaggag gaaatggeee accatgeect getqeqqaa
                                                                     240
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teetgggagg cageecagga gaeetgggag gaeaagegte gagggettae ettgeeceet
                                                                      300
ggcttcaaag cccagaatgg aatagccatt atggtctaca ccaactcatc gaacaccttg
                                                                      360
tactgggagt tgaatcangc cgtgcggacg ggcggaggct cccgggagct ctacatgagg
                                                                      420
caettteect teaaggeest geatttetae etgateeggg eestgeaget getg
                                                                      474
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     <211> 1222
     <212> DNA
     <213> Homo sapiens
     <400> 754
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gccggtgccc cccatgttgg aacctgagtt ggagattatc tcctaagcag atacctgctt
                                                                      120
ccaaactggg gatgtagggc ttggaaacta aaaaatgcca ggtctgaggg agaggaaaga
                                                                      180
acaagtecag caatacacag agetetgtgt atteagaggg aagttggeag ggttgtgtte
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gggcagagaa actccgagtg gtacaaaggg gacgtgcca gagtggagaa atcatgctaa
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ttgtctgcac tagagctgga gaacgccacc caaaatgaag agagaaaggg gagccctgtc
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cagageetee agggeeetge geettgetee ttttgtetae ettettetga tecagacaga
                                                                      420
ccccctggag ggggtgaaca tcaccagccc cgtgcgcctg atccatggca ccgtggggaa
                                                                      480
gteggetetg etttetgtge agtacageag taccageage gacaggeetg tagtgaagtg
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gcagctgaag cgggacaagc cagtgaccgt ggtgcagtcc attggcacag aggtcatcgg
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caccetgegg cetgactate gggacegtat cegactettt gaaaatgget ceetgettet
                                                                      660
cagegacetg cagetggeeg atgagggeac ctatgaggte gagateteea teacegaega
                                                                      720
                                                                      780
caccttcact ggggagaaga ccatcaacct tactgtagat gtgcccattt cgaggccaca
ggtgttgggg gcttcaacca ctgtgctgga gctcagcgag gccttcacct tgaactgctc
                                                                      840
                                                                      900
acatgagaat ggcaccaage ccagetacac ctggctgaag gatggcaage ccetectcaa
                                                                      960
tgactcgaga atgetectgt ecceegacea aaaggtgete accateacee gegtgeteat
ggaggatgac gacctgtaca gctgcgtggt ggaaaacccc atcaaccagg gccggaccct
                                                                     1020
geettgtaag ateacegaat acagaaaaag eteeetttea teaatttgge teeaggagge
                                                                     1080
attttcctcc ttgggacctt ggtgaagacc tggccaacaa gggaaaaccc cgtctttatt
                                                                     1140
aaaaatacaa aaaatgcccc cgctttgggt gtaagggcct gttttcccgc gcccttcggg
                                                                     1200
aggttttgaa cagtaaatct cc
                                                                     1222
     <210> 755
     <211> 667
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) ... (667)
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                                                                      120
aggtggctgg gaagaactct ccaacaataa atacatttga taagaaagat ggctttaaaa
                                                                      180
gtgctactag aacaagagaa aacgtttttc actcttttag tattactagg ctatttgtca
                                                                      240
tgtaaagtga cttgtgaatc aggagactgt agacagcaag aattcaggga tcggtctgga
                                                                      300
aactgtgttc cctgcaacca gtgtgggcca ggcatggagt tgtctaagga atgtggcttc
                                                                      360
ggctatgggg aggatgcaca gtgtgtgacg tgccggctgc acaggttcaa ggaggactgg
                                                                      420
ggettecaga aatgeaagee etgtetggae tgegeagtgg tgaacegett teagaaggea
                                                                      480
aattgttcag ccaccagtga tgccatctgc ggggactgct tgccaggatt ttataggaag
                                                                      540
```

```
acquaacttq tcqqctttca aqacatqqaq tqqtqqtnqq cccttqttqq qaqaacccc
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tteetteect ecetttaegg aaacceggca cttggttgec agccaagggt ccaaacette
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ggggaaa
                                                                       667
     <210> 756
     <211> 411
     <212> DNA
     <213> Homo sapiens
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     <221> misc feature
     <222> (1)...(411)
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                                                                       120
aaatccctga agaaacttaa atgtcctgct cctgtccgcc ctgcttcttc accctcttcc
                                                                       180
tecaetetat ttgccaagae ateteetggt tteatececa aaeteecaee ttagattete
                                                                       240
tottaaactg gatagatgat ctcatctttt acggcactct gtataacttc ttcccagaag
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agacgeetet gtttacette etaeteacte tatatetate eeteetgete etttggetae
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     <212> DNA
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caggaggagg ggaggagaga gtggggctct tctatcggaa ccccctcccc atgtggatcc
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gccccaagcg gaggtcgcgg aggaggttat cgaaaatatg cccgccctgc gccccgcttt
                                                                      240
gctgtgggcg ctgctgaqcc tatggctgtg ctgcgcgacc cccgcgcctg cattgcaatg
                                                                      300
                                                                      360
teetgaagge tatgaaceet eeccaetaga eegaaagtge geteeetaee eeaatgteag
                                                                      388
acgatectge ceatgeceag aaggtttt
     <210> 758
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     <212> DNA
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                                                                      120
atcagtaaac agcaacacaa caatcaactg ggcctttttg atgaagacaa aaccatagag
                                                                      180
gaaaaccatt agaagaggta ataaaggccc ttcttataca gttaatagag agcctcctgg
                                                                      240
atggaacaag accagctgtt gctactgaaa atttacttct gttttcaagt tcaaatagag
                                                                      300
actaaaacat tatcttcacg ggaattgatt ttacgtcttc caaacacata tgccacctta
                                                                      360
attgtgattt gtgtgatagt tcagctqctg aaagctttcg tttatctcta cctggttaaa
                                                                      420
```

```
caactttaaa taataacaag tcaatatatc tgtttattga ccagggttct tctcatcccc
                                                                      480
agagcacact gttgaagaag aaggtactta accetttgtt teectageee tgecacatat
                                                                      540
ctcatttttc acattctcaa tggggagata taattgttta aaaaatggaa tgaagccggg
                                                                      600
tggcatggct tacacttgta attccagcta tttgggaggc taaggcagga ggattgctcg
                                                                      660
gggcccggag ttcaagacca gtctaggcaa catagtgaga ccccatctct acaaaaaata
                                                                      720
aaaactaaca ccccgggttc ctgactactc aaaagggtga ggcagaggat cacttgagcc
                                                                      780
cagaagcaga agctgggtga gctagactgg gcacgcactc ctcatggtgc agaagaaacc
                                                                      840
tgc
                                                                      843
     <210> 759
     <211> 647
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                                                                      120
catgcccage ggetgccgct gcctgcatct cgtgtgcctg ttgtgcattc tgggggctcc
                                                                      180
eggteageet greegageeg argaergeag ereceaetgr gaeetggeee aeggergerg
                                                                      240
tgcacctgac ggctcctgca ggtgtgaccc gggctgggag gggctgcact gtgagcgctg
                                                                      300
tgtgaggatg cctggctgcc agcacggtac ctgccaccag ccatggcagt gcatctgcca
                                                                      360
cagtggctgg gcaggcaagt tctgtgacaa agatgaacat atctgtacca cgcagtcccc
                                                                      420
ctgccagaat ggaggccagt gcatgtatga cgggggcggt gagtaccatt qtqtqttt
                                                                      480
accaggette catgggegtg actgegageg caaggetgga eeetgtgaac aggeaggete
                                                                      540
cccatgccgc aatggcgggc agtgccagga cgaccagggc tttqctctca acttcacqtq
                                                                      600
ccgctgcttg gtgggctttg tgggtgcccg ctgtgacgtg taaggtg
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     <212> DNA
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     <400> 760
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                                                                      120
ggagagggac tctgtctcaa aaaaaaactg aggtcaggga gggtgagatg acggtgagag
                                                                      180
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gegecaagta cattgtetet gtgggetaec ageatgaeat gategteaac gtgtgggeet
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gaacagccaa gtggagggac cacggtattg cagcgtttgc tacaagagca gcttcgctat
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cagtcagaaa acctcatcat ggagaagcag ctgtctcctc gaatgcaaaa taatgaagaa
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ccacctgccc ccgcaaggcg gcatctgctg gtcctgctgc tgctcctctc taccctggtg
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acaggeetee agageetaet ceaaggette ageegaettt teetgaaagg taaeetgett
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cgggtggcct tctggatcat taagctgcca cggcggaggt cccaccagga tgccctggag
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                                                                     1020
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gcttctgttg ctqqtcttqc cttqqctcaq tcctqctaac tacattqaca atqtqqqcaa
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cctgcacttc ctqtattcaq aactctqtaa aqqtqcctcc cactacqqcc tqaccaaaqa
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taggaagagg cgctcacaag atggctgtcc agacggctgt gcgagcctca cagccacqqc
                                                                      420
tecetececa gaggtttetg cagetgecae cateteetta atgacagaeg ageetgeet
                                                                      480
agacaaccct gcctacgtgt cctcggcaga ggacgggcag ccagcaatca gcccagtgga
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atcatttaaa aaaataaatc gagctttgag tgttcttcga aggacaaaga gcgggagtgc
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     <211> 446
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accgccttgt gaatgtcttt ctgggcattc catttgccca gccgccactg ggccctgacc
                                                                      240
ggtteteage eecacaceca geacagecet gggagggtgt gegggatgee ageactgege
                                                                      300
ccccaatgtg cctacaagac gtggagagca tgaacagcag cagatttgtc ctcaacggaa
                                                                      360
aacagcagat etteteegtt teagaggaet geetggteet caaegtetat ageecagetg
                                                                      420
aggtccccgc agggtccggt aggccg
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     <210> 778
     <211> 416
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<212> DNA

<213> Homo sapiens

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     <222> (1) ... (416)
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                                                                      120
totateacce tggageetge ceageegage gaaggggaca acgteacget ggtegteeat
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gggetttegg gggaactget egectacage tggtatgegg ggeecacact cagegtgtea
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tacetggtgg ccagetacat cgtgagcaca ggcgatgaga ctcetggecc ggcccacacg
                                                                      300
gngcgggagg ctgtgcgccc cgatggcagc ctggacatcc agggcatcct gccccggcac
                                                                      360
tcaagcacct acatcctgca gaccttcaac aggcagttgc agaccgaggt gggctn
                                                                      416
     <210> 779
     <211> 382
     <212> DNA
     <213> Homo sapiens
     <400> 779
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aattttgcaa ccttgttctt ttgatgctag tttaacggat gaagagtccc ggaaaaattg
                                                                      180
ggaagaattt ggaaatccag atgggcctca aggtgtggta aatgatgatt ttaaaatatt
                                                                      240
ggcgatatgg tatatattat aaaaatgtta accagattaa aggaataata ttattttctt
                                                                      300
actaaactta tactcacatg gagtttaaca tagataaatt gagctetcat taatttttgc
                                                                      360
tttatttttc tttctaaaga cg
                                                                      382
     <210> 780
     <211> 437
     <212> DNA
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     <400> 780
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                                                                      120
                                                                      180
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aacagatgaa gatgcaagac gtgccataag tcgttcagga gggtttatca aggattcatc
                                                                      240
                                                                      300
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tgatcgtgta ggaagaggc gtccaggatc tgggacatca ggggttgaca gcctgtctaa
                                                                      360
ttttattgag tctgttaagg aagaagcaag taattctgga tatggctctt caattaatca
                                                                      420
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     <210> 781
     <211> 476
     <212> DNA
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<213> Homo sapiens

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                                                                      120
acatecectg tgggeacage eegegtgetg cagetggeet ttggetgeac tacetteage
                                                                      180
ctggtggctc accggggtgg ctttgcgggc gtccagggca ccttctgcat ggccgcctgg
                                                                      240
ggettetget tegeegtete tgegetggtg gtggeetgtg agtteacaeg geteeaegge
                                                                      300
tgcctgcggc tctcctgggg caacttcacc gccgccttcg ccatgctggc caccctgcta
                                                                      360
tgcgcgacgg ctgcggtcct gtatccgctg tactttgccc ggcgggagtg tccccccgag
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cocgoegget gtgctgccag ggacttecge ctggcagcca qtqtcttcqc cggqct
                                                                      476
     <210> 782
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     <212> DNA
     <213> Homo sapiens
     <400> 782
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                                                                      120
ataattttta tactcttctg catttgctaa atttcctctc attagcaggt tataccttta
                                                                      180
tgatcagaaa aaaaattaaa cactgcttct aaaaaatact catctccagc acttggagat
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cacctacctc tacattctac ccaactgagc ccaatttagt cttctcaggg ctttgcccaa
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gaacagttca ggaatgcatg cctctgaagg ccttcctgct cttccccttc tqqccttqqt
                                                                      360
atotoattot cattootgoo otococtaco totocaacco catcacttgo cagocatcot
                                                                      420
gttcttcctt gttggtcatc agttaatgaa gtgtattagg tgacctgagt acttgtcagt
                                                                      480
actteceaga ggeaagaaca tteetegeag ateaaggtae etttaagage caagaagete
                                                                      540
agatttggag gcgggagagc tgtactgcat cccctcaaat gttaqcagtg ccaaqaaatg
                                                                      600
agacgctagt ctagggggca ccacaagcag aaaggggctg tttcaaggag tcgtccgccc
                                                                      660
atgggagtct cctcttctat tattcacctt gctccaagga tatcttttct tttacqtatq
                                                                      720
aaaattttgt aattgttcaa ctataacacc atg
                                                                      753
     <210> 783
     <211> 769
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(769)
     <223> n = a,t,c or g
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                                                                      120
gataagtegg gettttggtg agacagaett teecaaceet etgeeeegee ggtgeecatg
                                                                      180
cttetgtgge tgetgetget gateetgaet eetggaagag aacaateagg ggtggeeea
                                                                      240
aaagctgtac ttctcctcga tcctccatgg tccacagcct tcaaaggaga aaaagtggct
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ctcatatgca gcagcatatc acattcccta gcccagggag acacatattg gtatcacgat
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gagaagttgt tgaaaataaa acatgacaag atccaaatta cagagcctgg aaattaccaa
                                                                      420
tgtaagaccc gaggatcctc cctcagtgat gccgtgcatg tggaattttc acctgactgg
                                                                      480
ctgatcctgc aggetttaca teetgttttt gaaggagaca atgtcattet gagatgtcag
                                                                      540
gggaaagaca acaaaaacac tcatcacaag gtttactaca aggatggaaa acagntttct
                                                                      600
aatagttata atttagagaa gaatacagtg gattcagtct cccgggataa tagcccatat
                                                                      660
tattgtgctg ggtaaaagag agtttacata cttgggattg gagaacttta aaacccccaa
                                                                      720
```

```
769
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     <210> 784
     <211> 979
     <212> DNA
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     <400> 784
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ccgtggcagt gaccagaagg ggccggaagg gggtggccgc cggccggggcc ccgccctggg
                                                                    240
geogecteec egegggttee gttggetgtg geggeagetg aegettgtgg eggeggtgge
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ttcggggtgg gcgtaagatg gcgacagcag cgcagggacc cctaagcttg ctgtggggct
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qqctqtqqaq cqaqcqcttc tqqctacccq agaacqtqaq ctqqqctqat ctqqaqqqqc
                                                                    420
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cgggcatett ettegtgagg etgetetteg agegatttat tgccaaacce tgtgcactee
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ccccaacqct tactaaattc tgtgaaagca tgtaagtacg caaggaggga gggagggaat
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     <212> DNA
     <213> Homo sapiens
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                                                                    120
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geatggeget cetgeteete caggegetge ceageceett gteagecagg getgaaceee
cgcaggataa ggaagcctgt gtgggtacca acaatcaaag ctacatctgt gacacaggac
                                                                    240
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tgtggaccat catcatcatc ctgagctgct gctgtgtttg ccaccaccgc cgagccaagc
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accgccttca ggcccagcag cggcaacatg aaatcaacct gatcgcttac cgagaagccc
                                                                    420
acaattactc agcgctgcca ttttatttca ggtttttgcc aaactattta ctacctcctt
                                                                    480
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                                                                    550
agcagcaacg
     <210> 786
     <211> 932
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) . . . (932)
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<223> n = a,t,c or g

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ccccttcatg ggcatcttgc cctccaacaa ggtctgcccc cagtacttcc tcgacaccaa
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gegaategee etcaagatee tgatqettqt tqqaqteeaq ecqqeeaqqe teateetqqq
                                                                      420
gatgatggtg accaectegt tettgteeat gtggetgage aacaecgeet ceaetgeeat
                                                                      480
gatgcttccc attgccaatg ccatcctgaa aagtctcttt ggccagaagg aggttcgaaa
                                                                      540
ggacccccag ccaggagat gaagagaaca cagggaatag aaccccaata cctntcctct
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ctgaggaaag gctgaaactt caagctcccc ttgtgataag acttggtcag ataactgagt
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ctggtcaatg gaatatgagt ggaaatgatg tgtgcaactt ccgggttctg tccttcctgc
                                                                      720
cgggtggaat gtgaatatga tggcacctgg gacccaaaga caggagccac atcttgagag
                                                                      780
atagatggca gatctgcccc tgtggctttg gatcatttac ctcagtgaac acaacaagca
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ttatccatga aaccataggt tttgtgtgct agttctagtt tttaaaatat gaattaaatt
                                                                      900
aaatacgtat ctgttaaaac ttaaaaaaaa aa
                                                                      932
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     <211> 514
     <212> DNA
     <213> Homo sapiens
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     <221> misc feature
     <222> (1)...(514)
     <223> n = a,t,c or g
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                                                                      120
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	gatgtgcccc					12780
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	cttgcccggg					13080
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	gctaaggctg					13500
	attacctctc					13560
	aatttatatg					13620
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	caggtagcaa					13860
	taggcatgtc					13920
	ccagggtgga					13920
	tgtatggccc					14040
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<211> 378

<212> DNA

<213> Homo sapiens

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     <213> Homo sapiens
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aaaaa
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cagtagtete tgaggageeg etegacette teeegaceet ggatetgagg caggagatge
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taaccgtccc cggttaatac cttttatcca tagccggcca ccacctcata cccatcccct
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gtggcctctg ctgagtctgc tgctcctgct gctgctgctc cagcctgtaa cctgtgccta
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caccacgcca ggcccccca gagccctcac cacgctgggc gcccccagag cccacaccat
                                                                      300
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tcagtctact ggtgccggga agactggcca aatcaggaaa tqaqqaaqat ctacaccact
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ccgan
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gaaatcagca tactggaaag ccctcaaggt gttcaagctg cctgtggagt tcctgctgct
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ggatttggtc tgtccatttt gaccaccatt ggtgagcaca tagtatacag gctgctgcta
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ccacgaatca aaaacaaatc caagctgcaa tactggctcc acaccagcca gagattacac
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agagcaataa atacatcatt tatagaggaa aagcagcagc atttcaagac caaacgtqtq
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gaaaagaggt ctaatgtggg accccgtcag cttaccgtat gg
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cctgtgtgga tgtgatatct agcagcatca ccggttactt acgttcgtat gtttttggtg
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gettettegt gtgcaatetg etgtatgege tgggeeecca cetgetggee tacegttgee
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tgcgggtggc ggatgaaatg gatgttaaca ttggtcatga ggttggctac gtgatccctt
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tegagaactg etgtaceaac gaaacaatec tgaggttggt ttgtggggtt cagteegete
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cctgctgatg attcttggct taggttctac aattctgaag gagcattatt ctqqcattct
                                                                    420
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cagttgcttt cccacact
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     <212> DNA
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atgagecage cacettgtgt etteceetee tatgacatag ceetteaget caceetacaa
                                                                    180
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                                                                    420
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                                                                    660
gtactacata gccagctgcc ttctcttcag acatctgcca gtactcatga gcagattctt
                                                                    720
acteccecgt gaaggetgte ttttgattgt ctttatgete tgtgaaaaga egetteettt
                                                                    780
cctgtttact ctaaaagaat acacatttat accagagcat aggacaactg atataaattg
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tgtaaacaca catgaaga
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    <210> 841
    <211> 459
    <212> DNA
    <213> Homo sapiens
    <220>
    <221> misc feature
    <222> (1)...(459)
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actactctag caaacctcat accetttact ctgagectaa tatgttttet getgttaate
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gtactcctgc tttgccaaac tgttgcaatc atgtatcctt cattccactc attcatcctg
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acqaataaat ctctqttqaa qaqataccat ttgacatttt agagatggct gcatgcaaac
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tcttaaaaca tttgaatgga ttttccctct tgttgcccag gctggagtgc aatggtgtga
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totoggttca otgcaaccco otgcotocog ggttcaagcg attotoctgo occagootoc
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tgagtagctg ggattagagg catgtgccac catgcccagc taattttgtg tttttagtag
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                                                                      424
ctgc
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     <213> Homo sapiens
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gacttttctc ttctgcatct atatcgattc gctcctctgt actgttccga agaacccagc
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caataatata atagaatggt acctttgttt atcgtctggt gtttttaaaa aatcaaacca
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atgatatact taaaaatacc tacttctgga tttgtaattt cagcaaagtt gaagatttag
                                                                      480
ctaacctaca ctatacccca gcttcactca ttgtccttaa catccaacag ttattagcca
                                                                      540
catcatgatt teetteagtt tatetaatgg ttgettttat aacttteaaa etatettett
                                                                      600
aaaatctatt tctggaacca tcacatttgg ctgggatcta agtaccaatg gaattccaat
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tgcaattaag aaccettaac ccactteett tttetta
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     <211> 698
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aggatcatgg ctgtaattag ggtcatggtg gtagttaggg tcacggctat agttggggtc
                                                                      240
atggtggtaa ttagggtcac agcgatagtt agcatcatgg tggtagttag ggtcatggtg
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gtagttaggg tcatggtggt agctaggccc atggtggtag ttagggtcat ggctgtagtt
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agagtcatgg cggatagtgc gctcagggct atatgttcgt cgtcgctgaa cgttacgttt
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tegettgaat agteaagece tgeetegtet titettitt teacteeaca aagaategte
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540

cttactcqaa tgcttttttc ccgtgcttaa ggtggcacac catccctggc caacatctct

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                                                                      600
eccgagttge gagageaatt etaaactage egttttageg tacceette actgaacetq
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aagegtettg teagateeea geacageeta eteeettggg eetgggeace teeagggetg
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agcggagggt acctggtggg gtgggctggg tcttactgca ggtgtgcctg gctcagggaa
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gagagetegt ggttggetgt geegttaeet tetteggatt gteagaetee agaetttggg
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ccagttctgc ccctcccagc acatgtgatg tgccagtgtg gtggactctt caagggagct
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gcctttaatc agagctggtt tgactta
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gagtgcctct ccaacccctg cagctctgag ggcagcctgg actgtataca gctcaccaat
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gatggtttca tttgccgttg tcccccggga ttttccqggg caaqqtqcca qaqcaqctqt
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ageccegtet etactcaaaa egeccaaaca attggeccag egttgtgggt ggeeteetet
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ggtcgccacc tacttcagag gtctgagcag cataactggt ttcgccccat atgccgtagg
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acgicgigga igcccitaat cggcticcic ggictiticac gcicaaggcc itagcccitic
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agaaatgcag accetgcgtg tggttttggc aaccetgggt gtgggagetg ettetettgg
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cctaagcata tattctcgac ccctgccctg gatcatctat ggagtctttg ccatcctctc
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gtttccgcaa gtacaaggtg gtgggtctct tcatgaagct gatggcgtcc atcatttcgg
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caggtaaaca ctatgtatta ggcaattaca gacctctaga gctattggtt ataaaagaag
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tetetacaaa aaatacaaaa attageeagg catagtgtca tgegeetgtg gteecageta
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420
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aattatggct cactgtagcc tcaccttcct gggatcaagc aatcttcttt cttcagcctc
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agacttaaga atgctggtga agagtgcaag agcctcaggg gccagcttga ggagcaaggc
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cggcagctgc aggctgctga ggaagctgtg gagaagctga aggccaccca agcagacatg
                                                                      240
ggagagaagt tgagctgcac tagcaaccat cttgcagagt gccaggcggc catgctgagg
                                                                      300
aaggacaagg agggggctgc cctgcgtgaa gaccaagaaa ggacccagaa ggaactcgaa
                                                                      360
aaaqccacqt gtattqcqqa cgaaatcqtc gacccqgqaa gtccggtccg aatgctqtca
                                                                      420
```

<210> 856

<211> 412

```
<212> DNA
     <213> Homo sapiens
     <400> 856
tttcgtcgcg ttctctcgct gcctgggctt ctgtggaatg agactcgggc tccttctact
                                                                       60
tgcaagacac tggtgcattg caggtgtgtt tccgcagaag tttgatggtg acagtgccta
                                                                      120
cgtggggatg agtgacggaa acccagaget cetgtcaacc agccagacet acaacqqcca
                                                                      180
gagegagaac aacgaagact atgagatece eccqataaca ceteccaace teccqqaqee
                                                                      240
atccctcctg cacctggggg accacgaagc cagctaccac tcgctgtgcc acggcctcac
                                                                      300
ccccaacggt ctgctccctg cctactccta tcaggccatg gacctcccag ccatcatggt
                                                                      360
gtccaacatg ctagcacagg acagccacct gctgtcgggc caqctqccca cq
                                                                      412
     <210> 857
     <211> 403
     <212> DNA
     <213> Homo sapiens
     <400> 857
eggteeggeg caaggaggge ggetggttgt ggaaaaagge etgggegage tgtgeetgea
                                                                       60
gcccctggct ggtttgggaa ggctgggctc ccaggctggt ggtagtggtg ggggtgattt
                                                                      120
tecteatgaa gececcaete egtecaetae tgeetgaeae eeaegaageg ageagtttee
                                                                      180
ggagetetee gatgtagggg cageaggtgt agageagetg etggteeace acaggeqeat
                                                                      240
tgtccaagcc atgctctggg gctactgtgt ccacctcaaa ggcatatgag ggaccctctt
                                                                      300
ccagaaagaa caagtcctca gggactgtgg gaatctggaa aagccagtcc agggcagcaa
                                                                      360
gaagcagcag cttgttcagg aaacacatct tcccctcact ctc
                                                                      403
     <210> 858
     <211> 439
     <212> DNA
     <213> Homo sapiens
     <400> 858
tgagggtggc gcaggggccc cggccagccc ggggctgcag cagtgcggac agctccagaa
                                                                       60
geteategge atetecattg geageetgeg egggetggge aceaagtgeg etgtgteeaa
                                                                      120
cgacctcacc gagcaggaga tacggacct ggagcattgt cccaattcct tcttctaatg
                                                                      180
aagaaatacg cttagttgat gatgcgtttg gaaaaatttg tcacatggtc agtgatgqct
                                                                      240
cttgggtggt tcgtgttcag gcaqcaaaac tqttgqqctc tatqqaqcaa qtcaqttctc
                                                                      300
atttettgga geagaceett gacaagaage atgteagate tgaggaggaa acqtaetqea
                                                                      360
catgagcgtg ccaaggaact ttacagttcg ggggagtttt ccagtggcag aaaqtgggga
                                                                      420
gatgatgctc ccaaggaag
                                                                      439
     <210> 859
     <211> 985
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) ... (985)
     <223> n = a,t,c or q
```

```
<400> 859
ggcagcatgg tggtgccgga gaaggagcag agctggatcc ccaagatctt caagaagaag
                                                                       60
acctgcacga cgttcatagt tgactccaca gatccgggga gcctggattg tcactggggg
                                                                      120
tetgeacaeg ggeateggee ggeatgttgg tgtggetgta egggaecate agatggeeag
                                                                      180
cactgggggc accaaggtgg tggccatggg tgtggccccc tgqggtgtgg tccggaatag
                                                                      240
agacaccete ateaacceca agggetegtt ceetgegagg taceggtgge geggtgacee
                                                                      300
ggaggacggg gtccagtttc ccctggacta caactactcg gccttcttcc tggtggacga
                                                                      360
cggcacacac ggctgcctgg ggggcgagaa ccgcttccgc ttgcgcctgg agtcctacat
                                                                      420
ctcacagcaa aacacggccg tggcagggac tggaattgac atccctggcc tgctcctcct
                                                                      480
gaaagaatgt gatgagaaga tggtgacgcg aatacacaac gccagccagg ctcagctccc
                                                                      540
atgtetteet tatgattgeg ttaaggggga getaeggaet tgeetagegg geaeceettg
                                                                      600
gaataccctc ttgcccccgg gaacggtggt tttccagcct acgccccgaa ccccgagaat
                                                                      660
gcatccacgc gcctcgtttt gctgaattga ngatccttgg acgtccttgc atcccacatc
                                                                      720
gtggcgaaat tatttatcta ccccccccg ccggtgggag taattgcata cttccatccc
                                                                      780
tattgcctcg ttttggagga gttggtgact ctcacttcta tcggtaatag gacattaccg
                                                                      840
tatecgacet tatgactegg tteccegate aacaategae tagtacegge egeggeeace
                                                                      900
tacctcctta taacacttct cttaccggca cctccgtcct tggtagtaaa ctcctggcgc
                                                                      960
tgtatctgtg tgctactgct aggcc
                                                                      985
     <210> 860
     <211> 396
     <212> DNA
     <213> Homo sapiens
     <400> 860
ctgcagaacc gagaggattc ttctgaaggc atcagaaaga agctggtgga agctgaggag
                                                                       60
ctcgaagaga aacatcggga ggcccaagtc tcagcccagc acctagaagt gcacctgaaa
                                                                      120
cagaaagagc agcactatga ggaaaagatt aaagtgttgg acaatcagat aaagaaagac
                                                                      180
ctggctgaca aggagacact ggagaacatg atgcagagac acgaggagga ggcccatgag
                                                                      240
aagggcaaaa ttctcagcga acagaaggcg atgatcaatg ctatggattc caagatcaga
                                                                      300
tecctggaac agaggattgt ggaactgtet gaagecaata aacttgcage aaatagcagt
                                                                      360
ctttttaccc aaaggaacat gaaggcccaa tgtatt
                                                                      396
     <210> 861
     <211> 686
     <212> DNA
     <213> Homo sapiens
     <400> 861
caagggaggg ctctgtgcca gccccgatga ggacgctgct gaccatcttg actgtgggat
                                                                       60
ccctggctgc tcacgcccct gaggacccct cggatctgct ccagcacgtg aaattccagt
                                                                      120
ccagcaactt tgaaaacatc ctgacgtggg acagcgggcc agagggcacc ccagacacgg
                                                                      180
tctacagcat cgagtataag acgtacggag agagggactg ggtggcaaag aagggctgtc
                                                                      240
ageggateae eeggaagtee tgeaacetga eggtggagae gggeaacete aeggagetet
                                                                      300
actatgccag ggtcaccgct gtcagtgcgg gaggccggtc agccaccaag atgactgaca
                                                                      360
ggttcagetc tetgcageac actaccetca agecacetga tgtgacetgt atetecaaag
                                                                      420
tgagatcgat tcagatgatt gttcatccta ccccacgcc aatccgtgca ggcgatgqcc
                                                                      480
accggctaac cctggaagac atcttccatg acctgttcta ccacttagag ctccaggtca
                                                                      540
accgcaccta ccaaatggtg agtgtatgtt gcaccctggt ctttctctgc ctaggaagcc
                                                                      600
tetteeetee caattagate tgagttgett taagaaaaaa aggggacatg ttatgtaaat
                                                                      660
tagcatttcc cacaacatgt cccttg
                                                                      686
```

```
<210> 862
     <211> 383
     <212> DNA
     <213> Homo sapiens
     <400> 862
cagagagttc aageccacac teeetgggeg tqtetqqetq qtqteaeett ttqqaqecaa
                                                                       60
cccctggtgg tggagtgtgg cagctgccct gcctgccctg ctgctgtcta tcctcatctt
                                                                      120
catggaccaa cagatcacag cagtcatcct caaccgcatg qaatacaqac tqcaqaaqqq
                                                                      180
agetggette cacetggace tettetgtgt ggetgtgetg atgetactea cateageget
                                                                      240
tggactgcct tggtatgtct cagccactgt catctccctg gctcacatgg acagtcttcg
                                                                      300
gagagagage agageetgtg eeceegggga gegeeeaae tteetgggta teaqqqaaca
                                                                      360
gaggctgaca ggcctggtgg tgt
                                                                      383
     <210> 863
     <211> 673
     <212> DNA
     <213> Homo sapiens
     <400> 863
caaccccaag accaagaagc acctgggcat tgccaaggtg gtctttgcca cggtccgggg
                                                                       60
agccaaggat gccgttcagc acttgcacag cacttccgtc atgggcaaca ttatccacgt
                                                                      120
ggagctggac accaaaggtg agcctggcag gggaggagcg tggggagacc tgtcagcccg
                                                                      180
accetttece tecceacet teetgeageg tggggaggae ceceeteae tetteettgg
                                                                      240
gatececece cacaacetta tttettagee eecteetgag ggtagagteg egtggageta
                                                                      300
aatgtgttgt ctgttgctag gagacagtct gtaatttacc aaatgtgccg gtccttggcc
                                                                     360
accgcacccc tagggaccac ccggaggctt ccccaccgct gacacccccg cgggccccct
                                                                      420
ctctgagccc tggtggcttg ggtttagaca gtccccagtg ttgcctgtgt taggggagga
                                                                      480
gacagagttt gtttacttgt gggggactga ggaagtgcca ctaggatgcc ttgaaataca
                                                                      540
tcaagagaag gtctgaaaac tgaaaagaga gtcctctaag gatccagggt gtcccccac
                                                                     600
ctccttgctg acccttcccc tctggaagtg gcagccaatc tggggcccag gaatgttqtt
                                                                     660
tcattgataa ggg
                                                                      673
     <210> 864
     <211> 435
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(435)
     <223> n = a,t,c or g
     <400> 864
gggaaatgtg tgggagccct gagcgtttgt gtgtgcgctg cgctcgtgtg tgcgctgtgt
                                                                      60
teatgegtge getgtgtgtt gtgtgtgtat atetgeggag acgeataaag tatgageget
                                                                     120
ttttaggatg ggaattgaga tgtaagattt gggggtgagg gccnccctga cccataggcc
                                                                     180
tgacatcctc atcctatgga ccctagagtc tggccactcc aggaacctga cctqctctqt
                                                                     240
geceegeeee tgtaageata gaacaceee catgatetee tggagtgggg ceteegagae
                                                                     300
```

```
ctccccgggc cccactactg cccgttcctc agtgctcacc cttaccccaa aqccccaqga
                                                                      360
nnaccggncc agccctcacc tgtnaggttg accttgcctg gggacagggt gtgacccacg
                                                                      420
accnatacct ntncg
                                                                      435
     <210> 865
     <211> 2161
     <212> DNA
     <213> Homo sapiens
     <400> 865
ggcggcgatg tcgctcgtgc tgctaagcct ggccgcgctg tgcaggagcg ccgtaccccg
                                                                       60
agageegace gtteaatgtg getetgaaac tgggeeatet ceagagtgga tgetacaaca
                                                                      120
tgatctaatc ccgggagact tgagggacct ccgagtagaa cctgttacaa ctagtgttgc
                                                                      180
aacaggggac tattcaattt tgatgaatgt aagctgggta ctccgggcag atgccagcat
                                                                      240
ccgcttgttg aaggccacca agatttgtgt gacgggcaaa agcaacttcc agtcctacag
                                                                      300
ctgtgtgagg tgcaattaca cagaggcctt ccagactcag accagaccct ctggtggtaa
                                                                      360
atggacattt tectacateg getteeetgt agagetgaae acagtetatt teattgggge
                                                                      420
ccataatatt cctaatgcaa atatgaatga agatggccct tccatgtctg tgaatttcac
                                                                      480
ctcaccaggc tgcctagacc acataatgaa atataaaaaa aagtgtgtca aggccggaag
                                                                      540
cctgtgggat ccgaacatca ctgcttgtaa gaagaatgag gagacagtag aagtgaactt
                                                                      600
cacaaccact cccctgggaa acagatacat ggctcttatc caacacagca ctatcatcgg
                                                                      660
gttttetcag gtgtttgage cacaccagaa gaaacaaacg cgagettcag tggtgattec
                                                                      720
agtgactggg gatagtgaag gtgctacggt gcagctgact ccatattttc ctacttgtgg
                                                                      780
cagegactge atcegacata aaggaacagt tgtgctctgc ccacaaacag gcgtcccttt
                                                                      840
ccctctggat aacaacaaaa gcaagccggg aggctggctg cctctcctcc tgctgtctct
                                                                      900
gctggtggcc acatgggtgc tggtggcagg gatctatcta atgtggaggc acgaaaggat
                                                                      960
caagaagact teetttteta eeaccacact actgeceece attaaggtte ttgtggttta
                                                                     1020
cccatctgaa atatgtttcc atcacacaat ttgttacttc actgaatttc ttcaaaacca
                                                                     1080
ttgcagaagt gaggtcatcc ttgaaaagtg gcagaaaaag aaaatagcag agatgggtcc
                                                                     1140
agtgcagtgg cttgccactc aaaagaaggc agcagacaaa gtcgtcttcc ttctttccaa
                                                                     1200
tgacgtcaac agtgtgtgcg atggtacctg tggcaagagc gagggcagtc ccagtgagaa
                                                                     1260
ctctcaagac ctcttccccc ttgcctttaa ccttttctgc agtgatctaa gaagccagat
                                                                     1320
tcatctgcac aaatacgtgg tggtctactt tagagagatt gatacaaaag acgattacaa
                                                                     1380
tgctctcagt gtctgcccca agtaccacct catgaaggat gccactgctt tctgtgcaga
                                                                     1440
actteteeat gteaageage aggtgteage aggaaaaaga teacaageet geeacgatgg
                                                                     1500
ctgctgctcc ttgtagccca cccatgagaa gcaagagacc ttaaaggctt cctatcccac
                                                                     1560
caattacagg gaaaaaacgt gtgatgatcc tgaagcttac tatgcagcct acaaacagcc
                                                                     1620
ttagtaatta aaacatttta taccaataaa attttcaaat attgctaact aatgtagcat
                                                                     1680
taactaacga ttggaaacta catttacaac ttcaaagctg ttttatacat agaaatcaat
                                                                     1740
tacagtttta attgaaaact ataaccattt tgataatgca acaataaagc atcttcagcc
                                                                     1800
aaacatctag tcttccatag accatgcatt gcagtgtacc cagaactgtt tagctaatat
                                                                     1860
totatgttta attaatgaat actaactota agaacccctc actgattcac tcaatagcat
                                                                     1920
cttaagtgaa aaaccttcta ttacatgcaa aaaatcattg tttttaagat aacaaaagta
                                                                     1980
gggaataaac aagctgaacc cacttttact ggaccaaatg atctattata tgtgtaacca
                                                                     2040
cttgtatgat ttggtatttg cataagacct tccctctaca aactagattc atatcttgat
                                                                     2100
tettgtacag gtgeetttta acatgaacaa caaaatacce acaaacttgt etaettttge
                                                                     2160
                                                                     2161
     <210> 866
     <211> 505
     <212> DNA
     <213> Homo sapiens
```

521

<220>

```
<221> misc feature
     <222> (1) ... (505)
     <223> n = a,t,c or q
     <400> 866
cataagcett gggcanagna cettgaaata aatgnggeea ceeacgegee cgeggaegeg
                                                                       60
tggggttgga atattctact ttgttattta tatcatcata tccttcctqq ttqtqqtqaa
                                                                      120
catgtacatt gcagtcatac tggagaattt tagtgttgcc actgaagaaa gtactgaacc
                                                                      180
totgagtgag gatgactttg agatgttcta tgaggtttgg gagaagtttg atcocqatqc
                                                                      240
gacccagttt atagagttct ctaaactctc tgattttgca gctgccctgg atcctcctct
                                                                      300
teteatagea aaacceaaca aagteeaget cattgeeatg gatetgeeca tggttagtgg
                                                                      360
tgaccggatc cattgtcttg acatcttatt tgcttttaca aagcgtgttt tgggtgagag
                                                                      420
tggggagatg gattetette gttcacagat ggaagaaagg ttcatgtetg caaateette
                                                                      480
caaagtgtcc tatgaaccca tcaca
                                                                      505
     <210> 867
     <211> 608
     <212> DNA
     <213> Homo sapiens
     <400> 867
ttcagttttt ggctctggtg caccatgtgc ctgggttaat ttgggtggct caatcccaaa
                                                                       60
gragetetga acceeaaage ggeteetetg aatteceagt tteaagttee actetgteee
                                                                      120
tgctgggcat ctcgagatat gggaaacagg gctgttataa ttgccagaca gctgagttct
                                                                      180
gtacatacct tgatttgcaa ttttttttgg ctgcttctca ggacaactgg gggagattta
                                                                      240
gattccttaa aatgcagtta tgaatctatt ggcctcaact ctatttctac ccatgaattc
                                                                      300
attigtacti ggcaaagacg acttaattic tcattigtta tgtcatttaa acctctcttt
                                                                      360
agageetete eteaetetta eetgttaata ateggaagte agetacatga aaegtteaat
                                                                      420
ttgggttcca tctcctctga agaaaaatgc agttaaaaaa aaaataagag gtttggccag
                                                                      480
ccgcagtggc tcacacctgt aatcccagca ttttgggagg ccgaggcagt cagatcacct
                                                                      540
gggggggga gttcgggaac cggcctggcc caacacagga gaaaccccgt cttatactaa
                                                                      600
acaatata
                                                                      608
     <210> 868
     <211> 772
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(772)
     <223> n = a,t,c or g
     <400> 868
tttcgtagcg caggcagggt tccctgctgg ggcccgggct gcccagccat gctttgggca
                                                                       60
ctctggccaa ggtggctggc agacaagatg ctgccctcc tgggggcagt gctgcttcag
                                                                      120
aagagagaga agaggggccc tctgtggagg cactggcggc gggaaaccta cccatactat
                                                                      180
gacctccagg tgaaggtgct gagggccaca aacatccggg gcacagacct gctgtccaaa
                                                                      240
gccgactgct atgtgcaact gtggctgccc acggcgtccc caagccctgc ccagactagg
                                                                      300
atagtggcca actgcagtga ccccgagtgg aatgagacct tccactacca gatccatggt
                                                                      360
gctgtgaaga acgtcctgga gctcaccctc tatgacaagg acatcctggg cagcgaccag
                                                                      420
etetetetge teetgtttga eetgagaage eteaagtgtg geeaacetea caaacacace
                                                                      480
```

```
540
ttcccactca accaccagga ttcacaagag ctgcaggtgg aatttgttct ggagaagagc
caggagectg catetgaagt cateaccaac ggggttetgg gggeteacce etggetgaga
                                                                      600
                                                                      660
atqaaqqqta tqattttqqq agaqqqqaqa gccccacggc aacagcacgg ccaatcttgg
gaggggggg tgggaccctc cccctctcc ccnngnanaa acaccggagg gaagatagtt
                                                                      720
qqqttttggg aaqaaatggc gaatgggacc ggcgccccac cccgcccccc ct
                                                                      772
     <210> 869
     <211> 704
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
     <222> (1)...(704)
     <223> n = a,t,c or g
     <400> 869
tttcqtqqca tqatqagcat gattaccagc ctcggccact ggctgctgca gggcttttcc
                                                                       60
tqaqccatqq tqtcttctqc cqtcaaaggg cgaccctaac tgcatcctgc tggagtcgag
                                                                      120
aaaaccaqqt aqactqqaaa qqatqtqtct acagtaactg aaacacatca ctgcgttttg
                                                                      180
ttacaqtcaa tqataqqqca qatctqaqtt ccaqagcacg gctcacagac ctttccttgc
                                                                      240
atcagtctgt gccgaagtcn nnnnnnnnc ttttttcttt ttttgcccac attacatcac
                                                                      300
ttcataattt accacctacg tagcatgact gtatatttgg aatcatttct tcacaagttt
                                                                      360
tagaccatat taaaggaaca ctggcagaac cctgtttgat ttccctttcg tctgttcccc
                                                                      420
tacattgccc tcctggcccc cttgaggaac tagatgagcg attagaactg gccagaggtc
                                                                      480
cttggaggaa caacagcgaa acagaagcat tagtagcatt gtcctcccca gtctaacact
                                                                      540
tgtcggaccc ctgatgagca gacttccctg tggggtgttc atatccccat gccccgctca
                                                                      600
gtgggcttca tgtctgagtc atatttgcct gctttccttt gaggtggtgg gcgccaaggt
                                                                      660
tgtgacaaat gcccggagtc ctggagctcg ctgttacggt tttg
                                                                      704
     <210> 870
     <211> 389
     <212> DNA
     <213> Homo sapiens
     <400> 870
tttcgtgagg ctttgttctt ttgttctttg tgatagatct aattgctgct cactctttgg
                                                                       60
gtetgtactg egtttatgag etgtgacaet egeegtgaag gtetgeaget teaeteetga
                                                                      120
accagegaga ggaggaacce accagaagga ggaaaaegeg gaacacatet gaatateaga
                                                                      180
aggaacaaac tocagacacg cogcotttaa gaactgtaac agtcaccgcg agggtccgtg
                                                                      240
gtttcattct tgaagtaagt gagaccaaga acctgccaat ttcagacaca atggagagcg
                                                                      300
ccagtcctgc tgcggggcca tacatctatt taatttcctc tcatcttccc cccggttccg
                                                                      360
                                                                      389
agaggaaggt gctttcacct gcactgttc
     <210> 871
     <211> 643
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
```

```
<222> (1) ... (643)
     <223> n = a,t,c or g
     <400> 871
tttcgtggat ggagccctcc tcctgatcct gtagtggtag taagaatcac cagcgcgggc
                                                                       60
aaggagtacg gacgggagtc agaggcagag cgagggtgtg tggagggccg gcggggaccg
                                                                      120
ccgggagcgc gcggatgtcg gtgttcctgg ggccagggat gccctctgca tctttattag
                                                                      180
taaatcttct ttcagcttta ctcatcctat ttgtgtttgg agaaacagaa ataagattta
                                                                      240
ctggacaaac tgaatttgtt gttaatgaaa caagtacaac agttattcgt cttatcattg
                                                                      300
aaaggatagg agagccagca aatgttactg caattgtatc gctgtatgga gaggacgctg
                                                                      360
gtgacttttt tgacacatat gctgcagctt ttatacctgc cggagaaaca aacagaacaq
                                                                      420
tgtacatagc agtatgtgat gatgacttac cagagcctga cgaaactttt atttttcact
                                                                      480
taacattaca gaaaccttca gcaaatgtga agcttggatg gccaaggact gttactgtga
                                                                      540
caatattatc aaatggacaa atggcatttt gggaatttat tttcatttta aatattggcc
                                                                      600
ttccccctcc aattccgcca agtggaagnt tgaaagcccc cct
                                                                      643
     <210> 872
     <211> 498
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1)...(498)
     <223> n = a,t,c or q
     <400> 872
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                                                                      120
agatgtactt gtgcttcctg ctggccgtgc tgctgcagct ctacgtggcc acggaggcca
                                                                      180
tecteattge actggttggg gecaegecat cetaccaetg ggacetggca gageteetge
                                                                      240
caaatcagag ccacggtaac cagtcagctg gtgaagacca ggcctttggg gactggctcc
                                                                      300
tgacagccaa cggcagtgag atccataagc acgtgcattt cagcagcagc ttcacctcta
                                                                      360
togoctogga gtggttttta attgccaaca gatcctacaa agtcagtgca gcaagctctt
                                                                      420
ttttcttcag tggtgtattt gttggagtta tctcttttgg tcagctttca gatcgcttcg
                                                                      480
gaaggaaaaa aqtctatc
                                                                      498
     <210> 873
     <211> 404
     <212> DNA
     <213> Homo sapiens
     <400> 873
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ggattecete eaggtgaega tgetetggtt eteeggegte ggggetetgg etgageqtta
                                                                     120
etgeegeege tegeetggga ttaegtgetg egtettgetg etaeteaatt getegggggt
                                                                     180
ccccatgtct ctggcttcct ccttcttgac aggttctgtt gcaaaatgtg aaaatgaagg
                                                                     240
tgaagtcctc cagattccat ttatcacaga caacccttgc ataatgtgtg tctgcttgaa
                                                                     300
caaggaagtg acatgtaaga gagagaagtg ccccgtgctg tcccgagact gtgccctggc
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catcaagcag aggggagcct gttgtgaaca gtgcaaaggt tgca
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     <211> 435
     <212> DNA
     <213> Homo sapiens
     <400> 874
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ataaqtqqcc ttggctacaq qatqtactgq ttcacaaact tcctatatga catgctcttt
                                                                      180
tacttggttt ccgtctgcct gtgtgttgcc gttattgtcg ccttccagtt aacagctttt
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                                                                      300
cttccatgga tgtacctgat gtccagaatc ttttccagtt cggacgtggc tttcatttcc
                                                                      360
tatgtctcac taaacttcat ctttggcctt tgtaccatgc tcataaccat tatgccccgg
                                                                      420
                                                                      435
ttgctagcca tcatc
     <210> 875
     <211> 703
     <212> DNA
     <213> Homo sapiens
     <400> 875
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                                                                      120
ccctcaactt ctccttcagc cataaatcag acatctggtc cctgggctgc atcattctgg
                                                                      180
acatgaccag ctgctccttc atggatggca cagaagccat gcatctgcgg aagtccctcc
                                                                      240
gccagagccc aggcagcctg aaggccgtcc tgaagacaat ggaggagaag cagatcccgg
                                                                      300
atgtggaaac cttcaggaat cttctgccct tgatgctcca gatcgacccc tcggatcgaa
                                                                      360
taacgataaa gtgagctcag ggtcggggtt tattttaacc tgtggattta tctttcaaca
                                                                      420
tctctccacc ctaatacaag cacagctagt tggctttgta acgcctcaaa gaactccatc
                                                                      480
acagatgccc tgattatccc tgcacagctg ggctttgccc agttctggct ctcccaaacc
                                                                      540
gtgctgcggc gagtaatccc gaatgtacgg tggagtgagc agactgaccc ccaggaggca
                                                                      600
caggaggegt agececagg acceacgaca ettttagggt tecagaaaaa agtttteatt
                                                                      660
                                                                      703
caacataaaa aaaaaaaaat tootaaagac aaaaaaaaaa aaa
     <210> 876
     <211> 429
     <212> DNA
     <213> Homo sapiens
     <400> 876
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                                                                      120
getggetget ggeactgtge etggeetgge tgtggaccca cetgacettg getgeeetge
                                                                      180
                                                                      240
agecteccae tgccaeagtg cttgtgcage agggcaectg egaggtgatt geggeteaec
gctgctgcaa ccggaaccgc atcgaggagc gctcccagac ggtgaaatgc tcctgttttt
                                                                      300
ctggccaggt ggccggcacc acgcgggcaa agccctcctg cgtggacgac ctgctcttgg
                                                                      360
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                                                                      420
catcgtcct
                                                                      429
```

```
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     <211> 1140
     <212> DNA
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     <400> 877
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                                                                      120
gggggcatgg gcccaggctt ccagtactag cctctctgat ctgcagagct ccaggacacc
                                                                      180
tggggtctgg aaggcagagg ctgaggacac cggcaaggac cccgttggac gtaactggtg
                                                                      240
cccctaccca atgtccaagc tggtcacctt actagctctt tgcaaaacag agaaattcct
                                                                      300
catccactcg cagcagccgt gtccgcaggg agctccagac tgccagaaag tcaaagtcat
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gtaccgcatg gcccacaagc cagtgtacca ggtcaagcag aaggtgctga cctctttggc
                                                                      420
ctggaggtgc tgccctggct acacgggccc caactgcgag caccacgatt ccatggcaat
                                                                      480
ccctgagcct gcagatcctg gtgacagcca ccaggaacct caggatggac cagtcagctt
                                                                      540
caaacctggc caccttgctg cagtgatcaa tgaggttgag gtgcaacagg aacagcagga
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acatetgetg ggagatetee agaatgatgt geacegggtg geagacagee tgeeaggeet
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gtggaaagcc ctgcctggta acctcacagc tgcagtgatg gaagcaaatc aaacagggca
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cgagttccct gatagatcct tggagcaggt gctgctaccc cacgtggaca ccttcctaca
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agtgcatttc agccccatct ggaggagctt taaccaaagc ctgcacagcc ttacccaggc
                                                                      840
cataagaaac ctgtctcttg acgtggaggc caaccgccag gccatctcca gagtccagga
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cagtgccgtg gccagggctg acttccagga gcttggtgcc aaatttgagg ccaaggtcca
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ggagaacact cagagagtgg gtcagetgcg acaggacgtg gaggaccgcc tgcacqccca
                                                                     1020
gcactttacc ctgcaccgct cgatctcaga gctccaagcc gatgtgqaca ccaaattqaa
                                                                     1080
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     <213> Homo sapiens
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tgcgagctct gcaagtatga gttcatcatg gagaccaagc tgaagccact gagaaaatgg
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gagaagttgc agatgacgtc cagcgagcgc aggaagatca tgtgctcagt gacattccac
                                                                      240
gtcattgcca tcacatgtgt ggtctggtcc ttgtatgtgc tcattgaccg tactgctgag
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gagatcaagc aggggcaggc aacaggaatc ctagaatggc ccttttggac taaattggtg
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gttgtggcca tcggcttcac cggaggactt ctttttatgt atgttcagtg taaagtgtat
                                                                      420
gtgcaattgt ggaagagact caaggcctat aatagagtga tctatgttca aaactgtcca
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gaaacaagca aaaagaatat ttttgaaaaa tctccactaa cagagcccaa ctttgaaaat
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aaacatggat atggaatctg tcattccgac acaaactctt cttgttgcac agagcctgaa
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gacactggag cagaaatcat tcacgtctga ttgtgtgcgg gttgtcattt tcctggacat
                                                                      660
ccatgaagag ctgaaggaaa ttgtttactg ccaattgtat acctttctta tgtcctttaa
                                                                      720
tagcatagac tggacaggtg actatttata gtggcttctc tttttctaaa ccctccttag
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tetectagaa aacetteetg tgggeeagge atgeetgggt cetgeetetq cetqqeaqet
                                                                      840
ctgtgggaaa gtggaagacc ccatgatgac atcatgggga gccagcagag ttcctgccca
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tggtcttgag ctgaatgaga gaataaaatg ccaatcccaa gggaagagga ggagcagggg
                                                                      960
tgcccaggcc ctgataccca gccgcctcca gcttgcagtg gtccccagcc tggagcagag
                                                                     1020
cattggggag tgtctaagcc atgacgagaa gattccctct gcatcacggc gaacccccag
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<210> 879

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1080

1139

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     <213> Homo sapiens
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aatatttgcc cacggcctcc caggcccagg cccatgccac ctgggccccg gcatctgttt
                                                                      180
qaggatetge caatgtgete ttaactgagg acgaaggaag aacacettte tatgagtett
                                                                      240
qcaaagatta cctccttcag gccacaaata tttgagtgca cactacgtgc caggcactgt
                                                                      300
                                                                      360
gcagggctgc aggcatagag acagaatgta atctatctgg gccttggacc ccatagggag
                                                                      420
aggggaccac teaggteeat actteetttg gaettgggge tttggeettg ggaggggegg
aggtggcgtg gcaagatgaa aaagacatce tgcccccatc cacttgggca gagettet
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     <210> 880
     <211> 546
     <212> DNA
     <213> Homo sapiens
     <400> 880
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                                                                       60
                                                                      120
gageteaggt getgeggtee teeettetge etgaaggagg catatggeea ggggeteege
ctgacactca cgaggcagta tatgcggatg atgggagtgc atccagtgat ccatttcctg
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gcctggttcc tggagaacat ggctgtgttg accataagca gtgctactct ggccatcgtt
                                                                      240
                                                                      300
ctgaaaacaa gtggcatctt tgcacacagc aataccttta ttgttttcct ctttctcttg
gattttggga tgtcagtcgt catgctgagc tacctcttga gtgcattttt cagccaagct
                                                                      360
aatacagcgg ccctttgtac cagcctggtg tacatgatca gctttctgcc ctacatagtt
                                                                      420
ctattggttc tacataacca attaagtttt gttaatcaga catttctgtg ccttctttcg
                                                                      480
                                                                      540
acaaccgcct ttggacaagg ggtatttttt attacattcc tggaaggaca agagacaggg
                                                                      546
attcac
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     <212> DNA
     <213> Homo sapiens
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tttaaaatct gatatattgg cataaaagta attgtacata tatatatgaa tgtgatttat
                                                                      180
tttcctttac atcttttgt tgtgtacagc agggcatata cttctcttgt cttggttgga
                                                                      240
tgcacaaatc tgtgtgcagt gctttttgcc cgttgcctag acgatcactt ggtttctctg
                                                                      300
aggatgtctg gttctcgtaa agagtttgat gtgaaacaga ttttgaaaat cagatggagg
                                                                      360
tggtttggtc atcaagcatc atctcctaat tctacagttg acagccagca gggagaattt
                                                                      420
tggaaccgag gacagactgg agcaaacggt gggagaaagt ttttagatcc atgtagccta
                                                                      480
caattgcctt tggcttcaat tggttaccga aggtccagcc aactggattt tcagaattca
                                                                      540
ccttcttggc caatggcatc cacctctgaa gtccctgcat ttgagtttac agcagaagat
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tgtggcggtg cacattggct ggatagacca gaagtggatg atggcactag tgaagaagaa
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aatgaatctg attccagttc atgcaggact tccaatagta gtcagacatt atcatcctqt
                                                                      720
catactatgg agccatgtac atcagatgaa tttttccaaq cccttaatca tqccqaqcaa
                                                                      780
acatttaaaa aaatggaaaa ctatttgaga cataaacagt tgtgtgatgt aattttagtc
                                                                      840
gctggtgatc gcagaattcc agctcacaga ttggtgctct cctctgtctc agactatttt
                                                                      900
gctggcatgt ttactaat
                                                                      918
     <210> 882
     <211> 604
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) ... (604)
     <223> n = a,t,c or q
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cttctaataa aagctctttg tattctggac gagtcatttt ctgtctggac tacattattt
                                                                      180
tcactctaag attgatccac atttttactg taagcagaaa cttaggaccc aagattataa
                                                                      240
tgctgcagag gatgctgatc gatgtgttct tettectgtt cetetttgcg gngtggatgg
                                                                      300
tggcctttgg cgtggccagg caagggatcc ttaggcagaa tgagcagcgc tggaggtgga
                                                                      360
tattccgttc ggtcatctac gagccctacc tggccatgtt cggccaggtg cccagtqacq
                                                                      420
tggatggtac cacgtatgac tttgcccact gcaccttcac tgggaatgag tccaagccac
                                                                      480
tgtgtgtgga gctggatgag cacaacctgc cccggttccc cgagtggatc accatcccc
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ttqq
                                                                      604
     <210> 883
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     <213> Homo sapiens
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cctacagggc ctgctggaga agaccaatgg gtgcatggga tgaccggcag cttccctcaa
                                                                      120
gtggcttccc agagactact aggagaactt ggtcctatcg ctgccccac ctggaaqctq
                                                                      180
gacttaagga tcccccaaag aacggggcaa ttagaaacct cccacccaqc gaagggataa
                                                                      240
getteteaae teagteeeae caetetteat egeaaceete tgagtetgea geagaaaeaa
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acatetecaa gttacagagg aggggatgga atececaagg ggeegagegg tageeetttt
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aacttataag cctgttgatt agcctatacg agttatttgc acgtcaagaa aggaagtagc
                                                                      420
etgeteette etgeagegte etgetggtgt gacageaegt eeccaagete agtgetaace
                                                                      480
toottattaa acateeeetg etgtgaetea gggaaceeae atgggtaete taaaacagte
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attcagggac cccacggggt catgtgggag ggagacagat cccagaaaga gcacaagtga
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aaggcaggtg aaagcaagcc gagccccact gctgaaggac aaagccacaq gaagcctqat
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gacatettte etetgagget tecaaaegat caceccaaat tgettgetga tactgggaag
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agtggccatg aactctccat tgctctgctg gctgtggaat gtttgctcag cacaggaagc
                                                                      900
atttaaggag aaagtcaaag tagccaaaag gcaaaccaga tggtggtgga catgtgggtg
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acagagcatc ctgcatttgt tgcctcgggg tgcagcccca aagataaagc cagcagtgtg
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```
1080
caaatgacaa atgetacccc aceteegeca ggeagecaga gecagggeeg aaggaegegg
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aaaqqaactq qtqtqqaaac ctgcccagga accgcactct caactgagaa gagtccgggg
                                                                     1200
cgcgtccccg cccggccgcc cggctgtgaa ttccgccaca cggcctaggg tgctcgaggt
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ctcgat
     <210> 884
     <211> 420
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
     <222> (1)...(420)
     <223> n = a,t,c or g
     <400> 884
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cggtgcagca gtattcgccc aacgggcgta tcggaaacca ctgatctcgt tccttgtcgg
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                                                                      240
cttctcgatg ctggcggccg gcgtaaccag tgcggcggga ctcgccctcg ccttctcggg
                                                                      300
cgactatete aaageettea tegacgteee aacegtteea geggegeteg tetteetget
                                                                      360
cctggtggga cttctcaatg ccagaggcat caaggagtcc atgcgcgcca ncgtcgtcat
gacagtegtg gaagteaceg ggetegteet egttgtegte etegegeteg tgeeaggeag
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     <211> 1696
     <212> DNA
     <213> Homo sapiens
     <400> 885
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                                                                      120
                                                                      180
gcagccaggc catggagctc tetgatgtca cecteattga gggtgtgggt aatgaggtga
tggtggtggc aggtgtggtg gtgctgattc tagccttggt cctagcttgg ctctctacct
                                                                      240
                                                                      300
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                                                                      540
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                                                                      780
geogeetget acaagaceea geoegeacae tgegttetet gaacattace gacaactgtg
tgattcactg ccaccgctca cccccagggt cagctgttcc aggcccctca gcctccttgg
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ggatgtatgg acgataagga cataggaaga aaatgaaagg gtcctctgaa ggagttcaaa
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gctgctggcc aagctcagtg gggagcctgg gctctgagat tccctcccac ctgtggttct
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gactettece agtgteetge atgtetgeec ceageaceca gggetgeetg caagggeage
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teageatgge eccageacaa etcegtaggg ageetggagt atcetteeat tteteageea
                                                                     1260
aatactcatc ttttgagact gaaatcacac tggcgggaat gaagattgtg ccagccttct
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cttatgggca cctagccgcc ttcaccttct tcctctaccc cttagcagga atagggtgtc
                                                                     1380
```

```
1440
gageteagte aggaaggga tggggeacea ageeaageee ceageattgg gageggeeag
                                                                    1500
gccacagctg ctgctcccgt agtcctcagg ctgtaagcaa gagacagcac tggcccttgg
                                                                    1560
ccagcgtcct accctgccca actccaagga ctgggtatgg attgctgggc cctaggctct
                                                                    1620
tgcttctggg gctattggag ggtcagtgtc tgtgactgaa taaagttcca ttttgtggtc
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ctgcaaaaaa aaaaaa
                                                                    1696
     <210> 886
     <211> 1410
     <212> DNA
     <213> Homo sapiens
     <400> 886
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                                                                    120
ggtgattatt tggctccgct catagccctg ccttcctcgg aggagccatc ggtgtcgcgt
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gcgtgtggag tatctgcaga catgactgcg tggaggagat tccagtcgct gctcctgctt
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ctcgggctgc tggtgctgtg cgcgaggctc ctcactgcag cgaagggtca gaactgtgga
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ggcttagtcc agggtcccaa tggcactatt gagagcccag ggtttcctca cgggtatccg
                                                                    360
aactatgcca actgcacctg gatcatcatc acgggcgagc gcaataggat acagttgtcc
                                                                    420
ttccatacct ttgctcttga agaagatttt gatattttat cagtttacga tggacagcct
                                                                    480
caacaaggga atttaaaagt gagattatcg ggatttcagc tgccctcctc tatagtgagt
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aactaatagc cctcttatgt ggtaaagagt tcatttttaa tgcagaagag tttcattaaa
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gctatgtaca gccataggat tgcctacaat gtttggttat attatttgtg gtgtacttct
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agataccage ttegtataaa eeattteaaa gatgteettt eaggtgteae gggaagtete
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tgaaccctca ggaagtcgct gtgcctgtta gtgaaggggc ggtgttactg gaa
                                                                    413
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```
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     <212> DNA
     <213> Homo sapiens
     <220>
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     <223> n = a,t,c or g
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ataggtcaag taagtaaata gagatttaaa aaattatgaa cacaaaggaa gtaacagcct
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tectgtettg etgtagtaac tgaccatatg egtttatate atgetaattg tgcaatttat
                                                                      240
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tggtttttgt agtaaaattc tcagttattt tttttcttcg ccaagataca gattaccttt
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                                                                      420
cctttaaget gatectaagg aagttatttt ttgtatacct tcagagaggg gataacatcc
                                                                      480
caaagatatt agtgttcaca gaggatggat atttcctacg agcctggaat tatacagttg
acacacctca tggtatattt gcagccagta ctctatatga acaatccgtc tggatcacgg
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                                                                      600
ttcaagtctt gggtactcca ggcaaaaaag gcactagttt gaatcctttg cagtttgata
                                                                      660
acccagcaga attatatgta gaggacacag gagatattta cattgtggat ggagatggag
                                                                      720
gattgaataa cagattgatc aaactgtccc aagatttcat gatcctttgg ctgcatggag
                                                                      780
aaaatgggac agggcctgct aagttcaaca tacctcacag tgttacactt gattcagctg
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     <213> Homo sapiens
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                                                                      240
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gttctaaggc ggggggcccg tgtccccaca gagcctggcc tggagccctg gaaggaggcc
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                                                                      420
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                                                                      480
                                                                      540
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cagtgtctgt ggaagagctg cgggaaggtg ctgagcacgg cgtcggcgat gcagagacac
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                                                                      660
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                                                                      840
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geetgeeage ggtteetgga etaagteegg etegtteaag aacataaget accacettet
                                                                     1140
                                                                     1200
coctcoccac cocctcoagg cocggggctg aaacagcccg aggacagccc caggggctgg
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cetececee gecaggtegg ggaggggtee caccacteaa agtgeeteta aagaaaceag
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gccaccctcc cgcctgtcgg cccgtagatt tatcaagggt gttatgggcc cagctttggg
                                                                     1440
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ctcacgtctg ggctgcacca ggcgaagaga gaaattaaag atttgaggtt tttccagaag
                                                                     1620
ctttgtctgc ctctcgggag gaaggccgtg gggctgggac cctgtggtgg gcaagtgggt
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ggagtetgge agetgeecae agagggeega gggteaeeeg teggeegeeg ceaeeceagg
                                                                     1740
cgaggccgga ggaaggatca tctgagacgc aggaggcatc tgctggagca gcaatttccc
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     <212> DNA
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gacttgtctt cggtagggac agtcaagtca ggcaaaaccg tgaacttggc tacagcaggc
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acaatcaagc cgggcacagc catgaatctg actacagttg ggacaaccaa gccagggatg
                                                                      240
gtcatggatt tgatagcctc agaaccagac aagctgggca aagccatggc tacaagaagc
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acagecaaac cagatatgac cacagagggt atagecatgg atteageaac ateagaccea
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gtcaagccgg acatgtatt
                                                                      379
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     <212> DNA
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ccatcctggc catctatgcc ggcgtcatca agtctgcctt cgaccccccg gacatcccgg
                                                                      180
tetgeeteet ggggaacege acgetgteae ggegeagett egatgeetge gteaaggeet
                                                                      240
acggcatcca caacaactca gccacctccg cgctctgggg cctcttctgc aacggctccc
                                                                      300
ageceagege egeetgtgae gagtaettea tecagaacaa egteacegaa atteagggea
                                                                      360
tcccgggcgc ggccagtggt gtcttcctgg agaaccg
                                                                      397
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     <212> DNA
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tgaatacatc tgggaggttg gtgtgggctt cgctcactcc ccccagccta actacatcca
                                                                      180
cgatatgaac cggatggagc tgctgaaact gctgctgaca tgcttctccg aggccatgta
                                                                      240
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cctgcccca gctccggaaa gtggcagcac caacccatgg gttcagttct tttgttccac
                                                                       300
                                                                       360
qqaqaacaqa catqccctqc ccctcttcac ctccctcctc aacaccgtgt gtgcctatga
-ccetgtggaa tacgggatee eetacaacca cctgtatt -
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agetggacaa gatgetggae eeccaggtgt ggegggagge agetacecag gtettetetg
                                                                       180
ccttgggcct gggctttggt ggtgtcattg ccttctccag ctacaataag caggacaaca
                                                                       240
actgccactt cgatgccgcc ctggtgtcct tcatcaactt cttcacgtca gtgttggcca
                                                                       300
ccctcgtggt gtttgctgtg ctgggcttca aggccaacat catgaatgag aagtgtgtgg
                                                                       360
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      <212> DNA
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 tctatttcta catgtggaag ttcgtgtctc ctctatgcat ggctgtgctc accacagcca
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 gcatcatcca gctgggggtc acgcccccgg gctacagcgc ctggatcaag gaggaggctg
                                                                       240
 ccgagcgcta cctgtatttc cccaactggg ccatggcacc cctgatcacc ctcatcgtcg
                                                                       300
 tggcgacgct gcccatccct gtggtgttcg tcctgcggca cttccaccta atctgtgatg
                                                                       360
                                                                       380
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      <211> 389
      <212> DNA
      <213> Homo sapiens
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      <221> misc feature
      <222> (1)...(389)
      <223> n = a,t,c or g
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                                                                       120
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                                                                       240
 tcaqcttqaq agacctgata gagatgatgt ctatcggcac gctcctggcc tacaccttgg
                                                                       300
 tetetgtetg tgtettgete ettegacace accetgagag tgacattgat ggttttgtea
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     <211> 415
     <212> DNA
     <213> Homo sapiens
     <400> 896
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acacctacat tctgttaaac aaactgggac ctgtgccctt tgaagggtta gaagagagcc
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caaatgggcc aaagatgggc ctcctgatga tgattctagg ccaaatattc ctgaatggca
                                                                      180
accaagccaa ggaggctgag atttgggaaa tgctctggag gatgggggtg cagcgggaaa
                                                                      240
ggaggettte catttttggg aacccaaaga gacttetgte tgtggagttt gtatggeage
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gttacttaga ctacaggcca gtaactgact gtaaaccagt ggagtatgag tttttctggg
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gcccaagatc ccacctagaa accaccaaga tgaaaattct gaagttcatg gcgaa
                                                                      415
     <210> 897
     <211> 428
     <212> DNA
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agaaagagtt tgttgctcag cccaactgcc aacagttgct tgccaccctg tggtatgatg
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getteeetgg atggeggegg aaacactggg tagtcaaget tetaacetge atgaccattg
                                                                      240
ggttcctgtt tcccatgctg tctatagcct acctgatctc acccaggagc aaccttgggc
                                                                      300
tgttcatcaa gaaacccttt atcaagttta tctqccacac aqcatcctat ttqaccttcc
                                                                      360
tetetatget teteetgget teteageaea ttgteaggae agaeetteat gtacaggge
                                                                      420
cctgtatt
                                                                      428
     <210> 898
     <211> 444
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc feature
     <222> (1) ... (444)
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caatgccatg ctacagttgg gccccttctt atattggaca tttctggctg cctttgaagg
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gacagtgttc ttctttggga cttactttct ttttcagact gcatccctag aagaaaatgg
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aaaggtatac ggaaactgga cttttggaac cattgttttt acagtcttaq tattcactqt
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aaccctgaag cttgccttgg atacccgatt ctggacgtgg ataaatcact ttgtgatttg
                                                                      360
gggttettta geettetatg tatttttete attettetgg ggaggaatta tttggeettt
                                                                      420
tctcaagcaa cagagaatgg cgaa
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```
<210> 899
     <211> 436
     <212> DNA
     <213> Homo sapiens
     <400> 899
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ageettteaa ggtgaacaga atgattteaa etecageeaa ggtgggaaag aettttgeea
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ccaacatggg ctgtttgagc accaaaaaac ccataatggg gagaggcctt atgagttcag
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tgaatgtggg gaattgttta ggtacaactc caaccttatt aaatatcagc aaaatcatgc
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tggagaaagg ccttatgagg gcactgaata tggaaagacc tttattagaa agtccaacct
                                                                      360
agttcagcac cagaaaattc acagtgaagg ctttctttca aaaaggtctg accccattga
                                                                      420
acatcaggag tgtatt
                                                                      436
     <210> 900
     <211> 466
     <212> DNA
     <213> Homo sapiens
     <220>
     <221> misc_feature
     <222> (1)...(466)
     <223> n = a,t,c or g
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ctgttgggcc tccgagagga ctgggatgac cgctggatca acgatgtgga agacagctac
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gggcagcagt ggacctatga gcagaggaaa atcgtggagt tcacctgcca cacagccttc
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ttcgtcagta tcgtgggggt gcagtgggcc gacttggtca tctgtaagac caggaggaat
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teggtettee ageeggggat gaagaacaag atettgatat ttggeetett tgaagagaca
                                                                      360
gccctggctg ctttcctttc ctactgccct ggaatgggtg ttgctcttaa gatgtatccc
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ctcaaaccta cctggagggt ctgtgccttc ccctactctc ttctca
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     <210> 901
     <211> 412
     <212> DNA
     <213> Homo sapiens
     <400> 901
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tgatgtctta ggagccccct ggaattggct gtacttcatc cccctcctca tcattggagc
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gagagtggag acccgaaggg ctttcatgaa gctgcggcgc cagcagcaga ttgagcgtga
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gctgaatggc taccgtgtct ggatagccaa agcagaggaa gtcatgctcg ctgaagaaaa
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                                                                      412
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gtattgcgca gcgatgcacg gccatcaagt accacttttc tcagcccatc cgcttgcgaa
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acatteettt taatttaace aagaceatae ageaagatga gtggeacetg etteatttaa
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gaagaatcac tgctggcttc ctcggcatgg ccgtagccgt ccttctctqc qqctqcattq
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tggccacagt cagtttcttc tgggaggaga gcttgaccca gcacgtggct ggactcctgt
                                                                      360
tecteatgae agggatattt tgeaceattt eeetetgtae ttatgeegee agtatetegt
                                                                      420
atgatttgaa ceggeteeca aagetaattt atageetgee tgetgatgtg gaacatqqtt
                                                                      480
acagetggte catettttge geetggtgea gtttaggett tattgtggea getggaggte
                                                                      540
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gccagccctt tctggattac tgatagaaaa tcatgcaaaa cctcccaacc tttctaaqqa
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aaattttaat geageatett cagaacttgt eetgatggtg tettattgtg teageaceaa
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gaaaagccca atcttcagct cccaaagtta ggaaaagtgt cagtagtcga atccatgaag
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ccgtgaaagc catcgtgctg tgtcacaacg tgacccccgt gtatgagtct cgggccggcg
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egetggteag cagggaeete acetecatge agetgaagae eeceagtgge caggteetea
                                                                      420
gettetgeat tetgeagetg ttteeettea eeteegagag caageggatg ggegteateg
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                                                                      600
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                                                                      600
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aaaaaaa
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1140

1200

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420

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<213> Homo sapiens

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<213> Homo sapiens

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Gly Gly Leu His Ser Ile Arg Thr Gly Met Arg Glu Arg Tyr His Ile
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Gln Gly Ser Val Gly His Asp Trp Ala Ala Leu Thr Phe Trp Leu Pro
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<210> 1011 <211> 83 <212> PRT <213> Homo sapiens

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<400> 1018 Met Leu Arg Phe Tyr Leu Ile Ala Gly Gly Ile Pro Leu Ile Ile Cys 70 Gly Ile Thr Ala Ala Val Asn Ile His Asn Tyr Arg Asp His Ser Pro 25 Tyr Cys Trp Leu Val Trp Arg Pro Ser Leu Gly Ala Phe Tyr Ile Pro 35 40 Val Ala Leu Ile Leu Leu Ile Thr Trp Ile Tyr Phe Leu Cys Ala Gly 55 Leu Arg Leu Arg Gly Pro Leu Ala Gln Asn Pro Lys Ala Gly Asn Ser 70 75 Arg Ala Ser Leu Glu Ala Gly Glu Glu Leu Arg Gly Ser Thr Arg Leu 90 85 Arg Gly Ser Gly Pro Leu Leu Ser Asp Ser Gly Ser Leu Leu Ala Thr 105 Gly Ser Ala Arg Val Gly Thr Pro Gly Pro Pro Glu Asp Gly Asp 120

<210> 1019 <211> 188 <212> PRT <213> Homo sapiens

<400> 1019 Met Gly Ser Ser Arg Leu Ala Ala Leu Leu Leu Pro Leu Leu Leu Ile 10 Val Ile Asp Leu Ser Asp Ser Ala Gly Ile Gly Phe Arg His Leu Pro 20 25 His Trp Asn Thr Arg Cys Pro Leu Ala Ser His Thr Asp Asp Ser Phe 40 Thr Gly Ser Ser Ala Tyr Ile Pro Cys Arg Thr Trp Trp Ala Leu Phe 55 Ser Thr Lys Pro Trp Cys Val Arg Val Trp His Cys Ser Arg Cys Leu 70 Cys Gln His Leu Leu Ser Gly Gly Ser Gly Leu Gln Arg Gly Leu Phe 85 90 His Leu Leu Val Gln Lys Ser Lys Lys Ser Ser Thr Phe Lys Phe Tyr 105 Arq Arq His Lys Met Pro Ala Pro Ala Gln Arg Lys Leu Pro Arg 125 120 Arg His Leu Ser Glu Lys Ser His His Ile Ser Ile Pro Ser Pro Asp 135 140 Ile Ser His Lys Gly Leu Arg Ser Lys Arg Thr Pro Pro Phe Gly Ser 150 155 Arg Asp Met Gly Lys Ala Phe Pro Lys Trp Asp Ser Pro Thr Pro Gly 170 165 Gly Asp Arg Pro Ser Ser Phe Glu Leu Leu Pro * 180 185 187

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<210> 1020
<211> 65
<212> PRT
<213> Homo sapiens
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<210> 1021 <211> 136 <212> PRT <213> Homo sapiens

<400> 1021 Met Pro Gly Phe Lys Phe Cys Ser Ser Leu Arg Phe Leu Tyr Leu Ile 10 Asn Phe Pro Ile Gly Lys Phe Val Cys Leu Ala Ile Leu Leu Pro His 20 25 Phe Pro Leu Leu Ser Cys Cys Pro Leu Gln Asp His Leu Asp Phe Pro 40 Gly Lys Glu Ser Arg Tyr Ser Gly Ser Cys Trp Leu Pro Ser Tyr Ser 55 60 Leu Ser Val Ala Gly Ser Pro Leu Gly His Leu Pro Asn Thr Tyr Met 70 75 His Thr Pro Arg Thr Phe Ser Leu Leu Pro Ile Pro His Pro Ser Val 85 90 Asn Trp Asp Ser Phe Lys Pro Phe Ser Ile Arg Glu Ala Leu Ala Thr 105 Val Glu Ser Leu Gly Arg Gln Ala Phe Pro Asn Thr Pro Thr Trp 120 Ala Phe Thr Leu His Leu Ser *

135

<210> 1022 <211> 186 <212> PRT <213> Homo sapiens

130

<400> 1022
Met Ala Gly Pro Arg Pro Arg Trp Arg Asp Gln Leu Leu Phe Met Ser

10 - 5 Ile Ile Val Leu Val Ile Val Val Ile Cys Leu Met Leu Tyr Ala Leu 25 30 20 Leu Trp Glu Ala Gly Asn Leu Thr Asp Leu Pro Asn Leu Arg Ile Gly 40 Phe Tyr Asn Phe Cys Leu Trp Asn Glu Asp Thr Ser Thr Leu Gln Cys 55 His Gln Phe Pro Glu Leu Glu Ala Leu Gly Val Pro Arg Val Gly Leu Gly Leu Ala Arg Leu Gly Val Tyr Gly Ser Leu Val Leu Thr Leu Phe Ala Pro Gln Pro Leu Leu Ala Gln Cys Asn Ser Asp Glu Arg Ala 100 105 Trp Arg Leu Ala Val Gly Phe Leu Ala Val Ser Ser Val Leu Leu Ala 125 120 Gly Gly Leu Gly Leu Phe Leu Ser Tyr Val Trp Lys Trp Val Arg Leu 135 140 Ser Leu Pro Gly Pro Gly Phe Leu Ala Leu Gly Ser Ala Gln Ala Leu 150 155 Leu Ile Leu Leu Ile Ala Met Ala Val Phe Pro Leu Arg Ala Glu 170 165 Arg Ala Glu Ser Lys Leu Glu Ser Cys * 180

<210> 1023 <211> 186 <212> PRT <213> Homo sapiens

Leu Trp Glu Ala Gly Asn Leu Thr Asp Leu Pro Asn Leu Arg Ile Gly
35 40 45
Phe Tyr Asn Phe Cys Leu Trp Asn Glu Asp Thr Ser Thr Leu Gln Cys

50 55 60

His Gln Phe Pro Glu Leu Glu Ala Leu Gly Val Pro Arg Val Gly Leu

65 70 75 80
Gly Leu Ala Arg Leu Gly Val Tyr Gly Ser Leu Val Leu Thr Leu Phe

85 90 95

Ala Pro Gln Pro Leu Leu Leu Ala Gln Cys Asn Ser Asp Glu Arg Ala
100 105 110

Trp Arg Leu Ala Val Gly Phe Leu Ala Val Ser Ser Val Leu Leu Ala 115 120 125

Gly Gly Leu Gly Leu Phe Leu Ser Tyr Val Trp Lys Trp Val Arg Leu 130 135 140

Ser Leu Pro Gly Pro Gly Phe Leu Ala Leu Gly Ser Ala Gln Ala Leu 145 150 155 160

Leu Ile Leu Leu Ile Ala Met Ala Val Phe Pro Leu Arg Ala Glu 165 170 175

Arg Ala Glu Ser Lys Leu Glu Ser Cys * 180 185

<210> 1024 <211> 73 <212> PRT <213> Homo sapiens

<400> 1024

 Met
 Val
 Cys
 Leu
 Val
 Gly
 Phe
 Leu
 Glu
 Leu
 Tyr
 Val
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 Phe
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 Ala
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 Met
 Glu
 Cys
 Asn
 Gly
 Thr

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 Thr
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 Asp
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 Pro

 35
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 Thr
 Arg
 His
 His
 Ala

 50
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 60
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<210> 1025 <211> 67 <212> PRT <213> Homo sapiens

<210> 1026 <211> 67 <212> PRT <213> Homo sapiens

<210> 1027 <211> 59 <212> PRT <213> Homo sapiens

<210> 1028 <211> 46 <212> PRT <213> Homo sapiens

<210> 1029 <211> 61 <212> PRT <213> Homo sapiens

<210> 1030 <211> 50 <212> PRT <213> Homo sapiens

<210> 1031 <211> 152 <212> PRT <213> Homo sapiens

<400> 1031 Met Ile Val Tyr Trp Val Leu Met Ser Asn Phe Leu Phe Asn Thr Gly 10 Lys Phe Ile Phe Asn Phe Ile His His Ile Asn Asp Thr Asp Thr Ile 20 25 Leu Ser Thr Asn Asn Ser Asn Pro Val Ile Cys Pro Ser Ala Gly Ser 40 Gly Gly His Pro Asp Asn Ser Ser Met Ile Phe Tyr Ala Asn Asp Thr 55 60 Gly Ala Gln Gln Phe Glu Lys Trp Trp Asp Lys Ser Arg Thr Val Pro 70 75 Phe Tyr Leu Val Gly Leu Leu Pro Leu Leu Asn Phe Lys Ser Pro 90 85 Ser Phe Phe Ser Lys Phe Asn Ile Leu Gly Ile Asn Asn Gln Val Ile 100 105 Leu Pro Gly Val Thr Glu Met Pro Gly Tyr Cys Pro Phe Leu Leu Pro 120 . 125 Val Ser Thr Glu Cys Cys Ala Val Ala Thr Ser Tyr Thr Cys Phe Glu 135 140 Glu Lys Asn Ile Gly Gln Cys Cys 150 152

<210> 1032 <211> 1764 <212> PRT <213> Homo sapiens

65					70					75					80
Leu	Thr	Glu	Met	Pro 85	His	His	Ser	Glu	Glu 90	Glu	Glu	Glu	Trp	Met 95	Ala
Gln	Ile	Leu	Gln 100	Ile	Leu	Thr	Val	Gln 105	Ala	Gln	Leu	Arg	Ala 110	Ser	Pro
		115	Pro				120					125			
	130		Met		_	135			_		140	-		_	
145			Gln		150					155					160
			Gly	165					170					175	
			Leu 180					185					190		
		195	Arg				200					205			
	210		Glu			215					220				
225			Asp		230					235					240
			Met	245					250					255	
			Val 260					265					270		
		275	Ser				280	_	_		_	285			
	290		Arg			295					300				
305			Leu		310		_			315					320
			Glu	325					330					335	
		_	340	_				345	_				350		
		355	Ala Ala			_	360					365	-		
	370		Leu			375					380				
385			Tyr		390					395	_				400
			Lys	405					410					415	
			420 Leu					425					430		
		435	Ala				440					445			
	450		Gly			455					460				
465		_	_		470		_			475					480
_	_	_	Ala	485	_	-			490					495	
	-	_	500	_				505	_				510		
		515	Asp				520	_				525			
атХ	530	ren	Phe	WIG	rne	535	мес	ьеи	cys	TIII	Met 540	nen	стА	ηλε	ьeu

Phe 545	Glu	Pro	Tyr	Val	Val 550	His	Val	Leu	Pro	His 555	Leu	Leu	Leu	Cys	Phe 560
Gly	Asp	Gly	Asn	Gln 565		Val	Arg	Glu	Ala 570		Asp	Asp	Cys	Ala 575	
Ala	Val	Met	Ser 580		Leu	Ser	Ala	His 585		Val	Lys	Leu	Val 590		Pro
Ser	Leu	Leu 595		Ala	Leu	Glu	Glu 600		Ser	Trp	Arg	Thr		Ala	Gly
Ser	Val 610		Leu	Leu	Gly	Ala 615		Ala	Tyr	Cys			Lys	Gln	Leu
Ser 625	Ser	Cys	Leu	Pro			Val	Pro	Lys		620 Thr	Glu	Val	Leu	
	Ser	His	Val		630 Val	Gln	Lys	Ala	_	635 Gln	Gln	Ala	Leu	_	640 Gln
Ile	Gly	Ser		645 Ile	Arg	Asn	Pro		650 Ile	Leu	Ala	Ile		655 Pro	Val
Leu	Leu		660 Ala	Leu	Thr	Asp		665 Ser	Arg	Lys	Thr		670 Lys	Cys	Leu
Gln	Thr	675 Leu	Leu	Asp	Thr		680 Phe	Val	His	Phe		685 Asp	Ala	Pro	Ser
	690 Ala	Leu	Ile	Met		695 Ile	Val	Gln	Arg		700 Phe	Gln	Asp	Arg	
705 Thr	Asp	Thr	Arg		710 Met	Ala	Ala	Gln		715 Ile	Gly	Asn	Met	_	720 Ser
Leu	Thr	Asp		725 Lys	Asp	Leu	Ala		730 Tyr	Leu	Pro	Ser		735 Thr	Pro
Gly	Leu		740 Ala	Ser	Leu	Leu		745 Pro	Val	Pro	Glu		750 Arg	Thr	Val
Ser	Ala	755 Lys	Ala	Leu	Gly		760 Met	Val	Lys	Gly		765 Gly	Glu	Ser	Cys
	770 Glu	Asp	Leu	Leu		775 Trp	Leu	Met	Glu		780 Leu	Thr	Tyr	Glu	
785 Ser	Ser	Val	Asp		790 Ser	Gly	Ala	Ala		795 Gly	Leu	Ala	Glu		800 Met
Ala	Gly	Leu		805 Val	Glu	Lys	Leu		810 Lys	Leu	Met	Pro		815 Ile	Val
Ala	Thr		820 Ser	Lys	Val	Asp		825 Ala	Pro	His	Val		830 Asp	Gly	Tyr
Ile	Met	835 Met	Phe	Asn	Tyr		840 Pro	Ile	Thr	Phe	Gly	845 Asp	Lys	Phe	Thr
	850 Tyr	Val	Gly				Pro	Cys				Ala	Leu	Ala	Asp
865 Glu	Asn	Glu	Phe		870 Arg		Thr	Ala		875 Arg		Gly	Gln		880 Val
Ile	Ser	Met	Tyr	885 Ala	Glu	Thr	Ala	Ile	890 Ala	Leu	Leu	Leu	Pro	895 Gln	Leu
Glu	Gln	Gly	900 Leu	Phe	Asp	Asp	Leu	905 Trp	Arg	Ile	Arg	Phe	910 Ser	Ser	Val
	Leu	915					920					925			
	930 Thr					935					940			_	-
945					950					955					960
	Asn			965					970					975	
	Leu		980					985					990		
	Gln	995				1	1000				1	L005			
Arg	Thr	Leu	Arg	Glu	Ile	Leu	Pro	Thr	Leu	Phe	Gly	Leu	Leu	Leu	Gly

1015 1020 Phe Leu Ala Ser Thr Cys Ala Asp Lys Arg Thr Ile Ala Ala Arg Thr 1030 1035 1040 Leu Gly Asp Leu Val Arg Lys Leu Gly Glu Lys Ile Leu Pro Glu Ile 1045 1050 1055 Ile Pro Ile Leu Glu Glu Gly Leu Arg Ser Gln Lys Ser Asp Glu Arg 1060 1065 1070 Gln Gly Val Cys Ile Gly Leu Ser Glu Ile Met Lys Ser Thr Ser Arg 1075 1080 1085 Asp Ala Val Leu Tyr Phe Ser Glu Ser Leu Val Pro Thr Ala Arg Lys 1090 1095 1100 Ala Leu Cys Asp Pro Leu Glu Glu Val Arg Glu Ala Ala Ala Lys Thr 1105 1110 1115 1120 Phe Glu Gln Leu His Ser Thr Ile Gly His Gln Ala Leu Glu Asp Ile 1125 1130 1135 Leu Pro Phe Leu Leu Lys Gln Leu Asp Asp Glu Glu Val Ser Glu Phe 1140 1145 1150 Ala Leu Asp Gly Leu Lys Gln Val Met Ala Ile Lys Ser Arg Val Val 1155 1160 1165 Leu Pro Tyr Leu Val Pro Lys Leu Thr Thr Pro Pro Val Asn Thr Arg 1170 1175 1180 Val Leu Ala Phe Leu Ser Ser Val Ala Gly Asp Ala Leu Thr Arg His 1185 . 1190 . 1195 . 1200 Leu Gly Val Ile Leu Pro Ala Val Met Leu Ala Leu Lys Glu Lys Leu 1205 1210 1215 Gly Thr Pro Asp Glu Gln Leu Glu Met Ala Asn Cys Gln Ala Val Ile 1220 1225 1230 Leu Ser Val Glu Asp Asp Thr Gly His Arg Ile Ile Glu Asp Leu 1235 1240 1245 Leu Glu Ala Thr Arg Ser Pro Glu Val Gly Met Arg Gln Ala Ala Ala 1250 1255 1260 Ile Ile Leu Asn Ile Tyr Cys Ser Arg Ser Lys Ala Asp Tyr Thr Ser 1270 1275 1280 His Leu Arg Ser Leu Val Ser Gly Leu Ile Arg Leu Phe Asn Asp Ser 1285 1290 1295 Ser Pro Val Val Leu Glu Glu Ser Trp Asp Ala Leu Asn Ala Ile Thr 1300 1305 1310 Lys Lys Leu Asp Ala Gly Asn Gln Leu Ala Leu Ile Glu Glu Leu His 1315 1320 Lys Glu Ile Arg Leu Ile Gly Asn Glu Ser Lys Gly Glu His Val Pro 1330 1335 1340 Gly Phe Cys Leu Pro Lys Lys Gly Val Thr Ser Ile Leu Pro Val Leu 1345 1350 1355 Arg Glu Gly Val Leu Thr Gly Ser Pro Glu Gln Lys Glu Glu Ala Ala 1365 1370 Lys Ala Leu Gly Leu Val Ile Arg Leu Thr Ser Ala Asp Ala Leu Arg 1380 1385 1390 Pro Ser Val Val Ser Ile Thr Gly Pro Leu Ile Arg Ile Leu Gly Asp 1395 1400 1405 Arg Phe Ser Trp Asn Val Lys Ala Ala Leu Leu Glu Thr Leu Ser Leu 1410 1415 1420 Leu Leu Ala Lys Val Gly Ile Ala Leu Lys Pro Phe Leu Pro Gln Leu 1430 1435 1440 Gln Thr Thr Phe Thr Lys Ala Leu Gln Asp Ser Asn Arg Gly Val Arg 1445 1450 1455 Leu Lys Ala Ala Asp Ala Leu Gly Lys Leu Ile Ser Ile His Ile Lys 1460 1465 1470 Val Asp Pro Leu Phe Thr Glu Leu Leu Asn Gly Ile Arg Ala Met Glu 1480

Asp Pro Gly Val Arg Asp Thr Met Leu Gln Ala Leu Arg Phe Val Ile 1490 1495 1500 Gln Gly Ala Gly Ala Lys Val Asp Ala Val Ile Arg Lys Asn Ile Val 1510 1515 Ser Leu Leu Ser Met Leu Gly His Asp Glu Asp Asn Thr Arg Ile 1525 1530 Ser Ser Ala Gly Cys Leu Gly Glu Leu Cys Ala Phe Leu Thr Glu Glu 1545 1550 Glu Leu Ser Ala Val Leu Gln Gln Cys Leu Leu Ala Asp Val Ser Gly 1560 1565 Ile Asp Trp Met Val Arg His Gly Arg Ser Leu Ala Leu Ser Val Ala 1575 1580 Val Asn Val Ala Pro Gly Arg Leu Cys Ala Gly Arg Tyr Ser Ser Asp 1590 1595 1600 Val Gln Glu Met Ile Leu Ser Ser Ala Thr Ala Asp Arg Ile Pro Ile 1605 1610 Ala Val Ser Gly Val Arg Gly Met Gly Phe Leu Met Arg His His Ile 1620 1625 1630 Glu Thr Gly Gly Gln Leu Pro Ala Lys Leu Ser Ser Leu Phe Val 1635 1640 1645 Lys Cys Leu Gln Asn Pro Ser Ser Asp Ile Arg Leu Val Ala Glu Lys 1655 1660 Met Ile Trp Trp Ala Asn Lys Asp Pro Leu Pro Pro Leu Asp Pro Gln 1670 1675 Ala Ile Lys Pro Ile Leu Lys Ala Leu Leu Asp Asn Thr Lys Asp Lys 1685 1690 Asn Thr Val Val Arg Ala Tyr Ser Asp Gln Ala Ile Val Asn Leu Leu 1700 1705 1710 Lys Met Arg Gln Gly Glu Glu Val Phe Gln Ser Leu Ser Lys Ile Leu 1720 1725 Asp Val Ala Ser Leu Glu Val Leu Asn Glu Val Asn Arg Arg Ser Leu 1735 1740 Lys Lys Leu Ala Ser Gln Ala Asp Ser Thr Glu Gln Val Asp Asp Thr 1750 1755 Ile Leu Thr * 1763

<210> 1033 <211> 151 <212> PRT <213> Homo sapiens

<400> 1033

 Met
 Asn
 Arg
 Ala
 Ser
 Gln
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 Leu
 Leu
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 Phe
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 Met
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 Jis

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 Ala
 Ile
 Ile
 Phe
 Val
 Pro
 Glu
 Glu
 Met
 Gln
 Met
 Leu
 Arg
 Arg
 Ala
 Ile
 Phe
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<210> 1034 <211> 149 <212> PRT <213> Homo sapiens

<400> 1034 Met Ala Leu Leu Pro Arg Trp Phe Arg Glu Ala Pro Val Leu Phe 1 5 10 Ser Thr Gly Trp Ser Pro Leu Asp Val Leu Leu His Ser Leu Leu Thr 25 Gln Pro Ile Phe Leu Ala Gly Leu Ser Gly Phe Leu Leu Glu Asn Thr 40 Ile Pro Gly Thr Gln Leu Glu Arg Gly Leu Gly Gln Gly Leu Pro Ser 55 Pro Phe Thr Ala Gln Glu Ala Arg Met Pro Gln Lys Pro Arg Glu Lys 70 75 Ala Ala Gln Val Tyr Arg Leu Pro Phe Pro Ile Gln Asn Leu Cys Pro 90 95 Cys Ile Pro Gln Pro Leu His Cys Leu Cys Pro Leu Pro Glu Asp Pro Gly Asp Glu Glu Gly Gly Ser Ser Glu Pro Glu Glu Met Ala Asp Leu 120 Leu Pro Gly Ser Gly Glu Pro Cys Pro Glu Ser Thr Arg Glu Gly Val Arg Ser Gln Lys * 145 148

<210> 1035 <211> 88 <212> PRT <213> Homo sapiens

<210> 1036 <211> 96 <212> PRT <213> Homo sapiens

<210> 1037 <211> 139 <212> PRT <213> Homo sapiens

<400> 1037 Met Ala Leu Ser Trp Met Thr Ile Val Val Pro Leu Leu Thr Phe Glu 10 Ile Leu Leu Val His Lys Leu Asp Gly His Asn Ala Phe Ser Cys Ile 25 Pro Ile Phe Val Pro Leu Trp Leu Ser Leu Ile Thr Leu Met Ala Thr 40 Thr Phe Gly Gln Lys Gly Gly Asn His Trp Trp Phe Gly Ile Arg Lys 55 Asp Phe Cys Gln Phe Leu Leu Glu Ile Phe Pro Phe Leu Arg Glu Tyr 70 75 Gly Asn Ile Ser Tyr Asp Leu His His Glu Asp Asn Glu Glu Thr Glu 90 Glu Thr Pro Val Pro Glu Pro Pro Lys Ile Ala Pro Met Phe Arg Lys 105 Lys Ala Arg Val Val Ile Thr Gln Ser Pro Gly Lys Tyr Val Leu Pro 120 125 Pro Pro Lys Leu Asn Ile Glu Met Pro Asp * 138 130 135

<210> 1038 <211> 64 <212> PRT <213> Homo sapiens

<210> 1039 <211> 286 <212> PRT <213> Homo sapiens

<400> 1039 Met Met Leu Gly Pro Val Thr Leu His Leu Val Gly His Leu Leu Ala 1 5 10 Phe Leu Asp Leu Cys Pro Arg Gly Pro Ile His Ser Ile Leu Pro 25 Met Thr Phe Glu Ala Val Lys Gln Asp His Gly Phe Met Leu Tyr Arg 40 Thr Tyr Met Thr His Thr Ile Phe Glu Pro Thr Pro Phe Trp Val Pro 55 Asn Asn Gly Val His Asp Arg Ala Tyr Val Met Val Asp Gly Val Phe 70 Gln Gly Val Val Glu Arg Asn Met Arg Asp Lys Leu Phe Leu Thr Gly 90 Lys Leu Gly Ser Lys Leu Asp Ile Leu Val Glu Asn Met Gly Arg Leu 105 Ser Phe Gly Ser Asn Ser Ser Asp Phe Lys Gly Leu Leu Lys Pro Pro 120 Ile Leu Gly Gln Thr Ile Leu Thr Gln Trp Met Met Phe Pro Leu Lys 135 Ile Asp Asn Leu Val Lys Trp Trp Phe Pro Leu Gln Leu Pro Lys Trp 150 155 Pro Tyr Pro Gln Ala Pro Ser Gly Pro Thr Phe Tyr Ser Lys Thr Phe 170 Pro Ile Leu Gly Ser Val Gly Asp Thr Phe Leu Tyr Leu Pro Gly Trp 1.8.5 180 Thr Lys Gly Gln Val Trp Ile Asn Gly Phe Asn Leu Gly Arg Tyr Trp 200 Thr Lys Gln Gly Pro Gln Gln Thr Leu Tyr Val Pro Arg Phe Leu Leu 215 220 Phe Pro Arg Gly Ala Leu Asn Lys Ile Thr Leu Leu Glu Leu Glu Asp 230 235 Val Pro Leu Gln Pro Gln Val Gln Phe Leu Asp Lys Pro Ile Leu Asn 245 250 Ser Thr Ser Thr Leu His Arg Thr His Ile Asn Ser Leu Ser Ala Asp 260 265 270 Thr Leu Ser Ala Ser Glu Pro Met Glu Leu Ser Gly His * 280

<210> 1040

<211> 96 <212> PRT <213> Homo sapiens

<400> 1040 et His Ala His

 Met His Ala His Ser Ala Ser Leu Trp Val Ala Phe Phe Tyr Arg Ser

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 10
 15

 Pro Phe Leu Phe Phe Thr Thr Gly Pro Pro Pro Pro Pro Thr Ser Ser Ser
 30

 Pro Ala Gly Leu Pro Leu Leu Glu Ser Thr Val Asp Ala Ser Arg Pro
 35
 40

 Asn Trp Leu Pro Leu Leu Leu Ser Pro Pro Leu Pro Phe Leu Ser Ile
 50
 60

 Glu Cys Thr Leu Tyr Asn Phe Ser Gly Ile Val Ile Glu Asn Lys Ile
 65
 70

 Phe Thr Ile Ile Thr Gly Phe Phe Gln Val Thr Ser Cys Arg Leu *
 85

<210> 1041 <211> 64 <212> PRT <213> Homo sapiens

<210> 1042 <211> 415 <212> PRT <213> Homo sapiens

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Asn Asn Cys Phe Ser Asp Ala Ile Val Val Cys Leu Thr Asn Cys Leu
                     120
Thr Ser Val Phe Ala Gly Phe Ala Ile Phe Ser Ile Leu Gly His Met
                 135
                                  140
Ala His Ile Ser Gly Lys Glu Val Ser Gln Val Val Lys Ser Gly Phe
                   155
               150
Asp Leu Ala Phe Ile Ala Tyr Pro Glu Ala Leu Ala Gln Leu Pro Gly
           165 170
Gly Pro Phe Trp Ser Ile Leu Phe Phe Phe Met Leu Leu Thr Leu Gly
             185
        180
Leu Asp Ser Gln Phe Ala Ser Ile Glu Thr Ile Thr Thr Ile Gln
         200
Asp Leu Phe Pro Lys Val Met Lys Lys Met Arg Val Pro Ile Thr Leu
                  215
Gly Cys Cys Leu Val Leu Phe Leu Leu Gly Leu Val Cys Val Thr Gln
    230
                               235
Ala Gly Ile Tyr Trp Val His Leu Ile Asp His Phe Cys Ala Gly Trp
                            250
            245
Gly Ile Leu Ile Ala Ala Ile Leu Glu Leu Val Gly Ile Ile Trp Ile
    , 260 265
Tyr Gly Gly Asn Arg Phe Ile Glu Asp Thr Glu Met Met Ile Gly Ala
         280 285
Lys Arg Trp Ile Phe Trp Leu Trp Trp Arg Ala Cys Trp Phe Val Ile
                295
                         300
Thr Pro Ile Leu Leu Ile Ala Ile Phe Ile Trp Ser Leu Val Gln Phe
             310
                      315
His Arg Pro Asn Tyr Gly Ala Ile Pro Tyr Pro Asp Trp Gly Val Ala
           325 330 335
Leu Gly Trp Cys Met Ile Val Phe Cys Ile Ile Trp Ile Pro Ile Met
            345
Ala Ile Ile Lys Ile Ile Gln Ala Lys Gly Asn Ile Phe Gln Arg Leu
      355 360
Ile Ser Cys Cys Arg Pro Ala Ser Asn Trp Gly Pro Tyr Leu Glu Gln
       375 380
His Arg Gly Glu Arg Tyr Lys Asp Met Val Asp Pro Lys Lys Glu Ala
              390
                              395
Asp His Glu Ile Pro Thr Val Ser Gly Ser Arg Lys Pro Glu *
                            410
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<210> 1043 <211> 48 <212> PRT <213> Homo sapiens

<210> 1044

<211> 146 <212> PRT <213> Homo sapiens

<400> 1044 Met Leu Phe Ser Ser Met Thr Leu Arg Leu Ser Arg Cys Ser Cys Ser 10 Ile Leu Leu Phe Trp Ala Ser Ala Ala Cys Met Phe Pro Ser Ser Arg 20 25 Tyr Leu Trp Ser Gly Arg Ser Leu Val Ser Val Glu Gly Ser Asp Arg 40 Phe Ser Ser Ala Val Ser Ser Phe Ser Ser Lys Ala Asn Trp Val Lys 55 Pro Lys Phe Arg Ser Trp Ser Gly Gly Ile Glu Leu Gly Phe Gln Met 70 His Trp Pro Pro Gly Val Gly Pro Arg Tyr Ser Pro Ser Cys His Phe 90 Pro Lys Ser Arg Trp Arg Thr Arg Pro Leu Arg Leu Ser Thr Ala Pro 105 Cys Thr Ser Trp Thr Leu Glu Leu Gln Tyr Leu Ala Leu Gln Lys Val 120 Ile Leu Gln Trp Gln Glu Leu Ser Cys Val Phe Arg Met Ser Thr Ser 135 Pro * 145

<210> 1045 <211> 53 <212> PRT <213> Homo sapiens

<210> 1046 <211> 407 <212> PRT <213> Homo sapiens

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40
Ser Arg His Ala Ala Glu Leu Arg Asp Phe Lys Asn Lys Met Leu Pro
                       55
Leu Leu Glu Val Ala Glu Lys Glu Arg Glu Ala Leu Arg Thr Glu Ala
Asp Thr Ile Ser Gly Arg Val Asp Arg Leu Glu Arg Glu Val Asp Tyr
                                  90
Leu Glu Thr Gln Asn Pro Ala Leu Pro Cys Val Glu Phe Asp Glu Lys
                            105
Val Thr Gly Gly Pro Gly Thr Lys Gly Lys Gly Arg Arg Asn Glu Lys
                         120
                                            125
Tyr Asp Met Val Thr Asp Cys Gly Tyr Thr Ile Ser Gln Val Arg Ser
                     135
                                        140
Met Lys Ile Leu Lys Arg Phe Gly Gly Pro Ala Gly Leu Trp Thr Lys
                  150
                              155
Asp Pro Leu Gly Gln Thr Glu Lys Ile Tyr Val Leu Asp Gly Thr Gln
              165
                    170 175
Asn Asp Thr Ala Phe Val Phe Pro Arg Leu Arg Asp Phe Thr Leu Ala
                   185 190
Met Ala Ala Arg Lys Ala Ser Arg Val Arg Val Pro Phe Pro Trp Val
                         200
Gly Thr Gly Gln Leu Val Tyr Gly Gly Phe Leu Tyr Phe Ala Arg Arg
                     215
Pro Pro Gly Arg Pro Gly Gly Gly Glu Met Glu Asn Thr Leu Gln
                  230
                                    235
Leu Ile Lys Phe His Leu Ala Asn Arg Thr Val Val Asp Ser Ser Val
              245
                                250
Phe Pro Ala Glu Gly Leu Ile Pro Pro Tyr Gly Leu Thr Ala Asp Thr
                             265
Tyr Ile Asp Leu Ala Ala Asp Glu Glu Gly Leu Trp Ala Val Tyr Ala
                         280
Thr Arg Glu Asp Asp Arg His Leu Cys Leu Ala Lys Leu Asp Pro Gln
                     295
Thr Leu Asp Thr Glu Gln Gln Trp Asp Thr Pro Cys Pro Arg Glu Asn
                  310
                                    315
Ala Glu Ala Ala Phe Val Ile Cys Gly Thr Leu Tyr Val Val Tyr Asn
                                 330
Thr Arg Pro Ala Ser Arg Ala Arg Ile Gln Cys Ser Phe Asp Ala Ser
                             345
Gly Thr Leu Thr Pro Glu Arg Ala Ala Leu Pro Tyr Phe Pro Arg Arg
                         360
                                            365
Tyr Gly Ala His Ala Ser Leu Arg Tyr Asn Pro Arg Glu Arg Gln Leu
                     375
                                        380
Tyr Ala Trp Asp Asp Gly Tyr Gln Ile Val Tyr Lys Leu Glu Met Arg
                                     395
                  390
Lys Lys Glu Glu Glu Val
              405 406
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<210> 1047

<211> 268

<212> PRT

<213> Homo sapiens

<400> 1047

Met Ile Gln Lys Ile Leu Phe Lys Asp Leu Phe Arg Phe Leu Leu Val 1 5 10 15

Tyr Leu Leu Phe Met Ile Gly Tyr Ala Ser Ala Leu Val Ser Leu Leu 20 Asn Pro Cys Ala Asn Met Lys Val Cys Asn Glu Asp Gln Thr Asn Cys 40 Thr Val Pro Thr Tyr Pro Ser Cys Arg Asp Ser Glu Thr Phe Ser Thr 55 Phe Leu Leu Asp Leu Phe Lys Leu Thr Ile Gly Met Gly Asp Leu Glu 70 Met Leu Ser Ser Thr Lys Tyr Pro Val Val Phe Ile Ile Leu Leu Val 85 90 Thr Tyr Ile Ile Leu Thr Phe Val Leu Leu Leu Asn Met Leu Ile Ala 105 Leu Met Gly Glu Thr Val Gly Gln Val Ser Lys Glu Ser Lys His Ile 120 Trp Lys Leu Gln Trp Ala Thr Thr Ile Leu Asp Ile Glu Arg Ser Phe 135 Pro Val Phe Leu Arg Lys Ala Phe Arg Ser Gly Glu Met Val Thr Val 150 155 -Gly Lys Ser Ser Asp Gly Thr Pro Asp Arg Trp Cys Phe Arg Val 165 170 Asp Glu Val Asn Trp Ser His Trp Asn Gln Asn Leu Gly Ile Ile Asn 180 185 Glu Asp Pro Gly Lys Asn Glu Thr Tyr Gln Tyr Tyr Gly Phe Ser His 200 Thr Val Gly Arg Leu Arg Arg Asp Arg Trp Ser Ser Val Val Pro Arg 215 220 Val Val Glu Leu Asn Lys Asn Ser Asn Pro Asp Glu Val Val Pro 230 235 Leu Asp Ser Met Gly Asn Pro Arg Cys Asp Gly His Gln Gln Gly Tyr 245 250 Pro Arg Lys Trp Arg Thr Asp Asp Ala Pro Leu * 260 265 267

<210> 1048 <211> 59 <212> PRT <213> Homo sapiens

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<210> 1049 <211> 77 <212> PRT <213> Homo sapiens

<210> 1050 <211> 474 <212> PRT <213> Homo sapiens

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<400> 1050 Met Arg Ala Leu Val Leu Leu Gly Cys Leu Leu Ala Ser Leu Leu Phe 1 5 Ser Gly Gln Ala Glu Glu Thr Glu Asp Ala Asn Glu Glu Ala Pro Leu Arg Asp Arg Ser His Ile Glu Lys Thr Leu Met Leu Asn Glu Asp Lys Pro Ser Asp Asp Tyr Ser Ala Val Leu Gln Arg Leu Arg Lys Ile Tyr 55 His Ser Ser Ile Lys Pro Leu Glu Gln Ser Tyr Lys Tyr Asn Glu Leu 70 Arg Gln His Glu Ile Thr Asp Gly Glu Ile Thr Ser Lys Pro Met Val 85 Leu Phe Leu Gly Pro Trp Ser Val Gly Lys Ser Thr Met Ile Asn Tyr 105 Leu Leu Gly Leu Glu Asn Thr Arg Tyr Gln Leu Tyr Thr Gly Ala Glu 120 Pro Thr Thr Ser Glu Phe Thr Val Leu Met His Gly Pro Lys Leu Lys 135 140 Thr Ile Glu Gly Ile Val Met Ala Ala Asp Ser Ala Arg Ser Phe Ser 150 155 Pro Leu Glu Lys Phe Gly Gln Asn Phe Leu Glu Lys Leu Ile Gly Ile 170 Glu Val Pro His Lys Leu Leu Glu Arg Val Thr Phe Val Asp Thr Pro 185 Gly Ile Ile Glu Asn Arg Lys Gln Gln Glu Arg Gly Tyr Pro Phe Asn 200 205 Asp Val Cys Gln Trp Phe Ile Asp Arg Ala Asp Leu Ile Phe Val Val 215 220 Phe Asp Pro Thr Lys Leu Asp Val Gly Leu Glu Leu Glu Met Leu Phe 230 235 Arg Gln Leu Lys Gly Arg Glu Ser Gln Ile Arg Ile Ile Leu Asn Lys 245 250 Ala Asp Asn Leu Ala Thr Gln Met Leu Met Arg Val Tyr Gly Ala Leu 265 270 Phe Trp Ser Leu Ala Pro Leu Ile Asn Val Thr Glu Pro Pro Arg Val 280 285 Tyr Val Ser Ser Phe Trp Pro Gln Glu Tyr Lys Pro Asp Thr His Gln 290 295

Glu Leu Phe Leu Gln Glu Glu Ile Ser Leu Leu Glu Asp Leu Asn Gln 305 . 310 . 315 . 320 Val Ile Glu Asn Arg Leu Glu Asn Lys Ile Ala Phe Ile Arg Gln His 330 Ala Ile Arg Val Arg Ile His Ala Leu Leu Val Asp Arg Tyr Leu Gln 340 345 Thr Tyr Lys Asp Lys Met Thr Phe Phe Ser Asp Gly Glu Leu Val Phe 355 360 365 Lys Asp Ile Val Glu Asp Pro Asp Lys Phe Tyr Ile Phe Lys Thr Ile 375 380 Leu Ala Lys Thr Asn Val Ser Lys Phe Asp Leu Pro Asn Arg Glu Ala 390 395 Tyr Lys Asp Phe Phe Gly Ile Asn Pro Ile Ser Ser Phe Lys Leu Leu 405 410 Ser Gln Gln Cys Ser Tyr Met Gly Gly Cys Phe Leu Glu Lys Ile Glu 425 Arg Ala Ile Thr Gln Glu Leu Pro Gly Leu Leu Gly Ser Leu Gly Leu 440 Gly Lys Asn Pro Gly Ala Leu Asn Cys Asp Lys Thr Gly Cys Ser Glu 455 Thr Pro Lys Asn Arg Tyr Arg Lys His * 470

<210> 1051 <211> 47 <212> PRT <213> Homo sapiens

<210> 1052 <211> 233 <212> PRT <213> Homo sapiens

85 90 Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Cys Ser Tyr Ala Gly Arg 100 105 110 Thr Thr Trp Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln 120 Pro Lys Ala Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser Glu Glu 135 Leu Gln Ala Asn Lys Ala Thr Leu Val Cys Leu Ile Ser Asp Phe Tyr 150 155 Pro Gly Ala Val Thr Val Ala Trp Lys Ala Asp Ser Ser Pro Val Lys 165 170 175 Ala Gly Val Glu Thr Thr Pro Ser Lys Gln Ser Asn Asn Lys Tyr 180 185 190 Ala Ala Ser Ser Tyr Leu Ser Leu Thr Pro Glu Gln Trp Lys Ser His 195 200 Arg Ser Tyr Ser Cys Gln Val Thr His Glu Gly Ser Thr Val Glu Lys 210 215 Thr Val Ala Pro Thr Glu Cys Ser * 230 232

<210> 1053 <211> 147 <212> PRT

<213> Homo sapiens

<400> 1053

Met Gly Ala Asp Arg Gly Pro His Val Val Leu Trp Thr Leu Ile Cys 10 Leu Pro Val Val Phe Ile Leu Ser Phe Val Val Ser Phe Tyr Tyr Gly 25 Thr Ile Thr Trp Tyr Asn Ile Phe Leu Val Tyr Asn Glu Glu Arg Thr 40 Phe Trp His Lys Ile Ser Tyr Cys Pro Cys Leu Val Leu Phe Tyr Pro 55 Val Leu Ile Met Ala Met Ala Ser Ser Leu Gly Leu Tyr Ala Ala Val 70 Val Gln Leu Ser Trp Ser Trp Glu Ala Trp Trp Gln Ala Ala Arg Asp 90 85 Met Glu Lys Gly Phe Cys Gly Trp Leu Cys Ser Lys Leu Gly Leu Glu 100 105 Asp Cys Ser Pro Tyr Ser Ile Val Glu Leu Leu Glu Ser Asp Asn Ile 120 125 Ser Ser Thr Leu Ser Asn Lys Asp Pro Ile Gln Glu Val Glu Thr Ser 135 140 Thr Val * 145 146

<210> 1054

<211> 123

<212> PRT

<213> Homo sapiens

<400> 1054

Met Tyr Val Thr Leu Val Phe Arg Val Lys Gly Ser Arg Leu Val Lys Pro Ser Leu Cys Leu Ala Leu Leu Cys Pro Ala Phe Leu Val Gly Val Val Arg Val Ala Glu Tyr Arg Asn His Trp Ser Asp Val Leu Ala Gly 40 Phe Leu Thr Gly Ala Ala Ile Ala Thr Phe Leu Val Thr Cys Val Val 55 His Asn Phe Gln Ser Arg Pro Pro Ser Gly Arg Arg Leu Ser Pro Trp 70 75 Glu Asp Leu Gly Gln Ala Pro Thr Met Asp Ser Pro Leu Glu Lys Asn 85 90 Pro Arg Ser Ala Gly Arg Ile Arg His Arg His Gly Ser Pro His Pro 105 Ser Arg Arg Thr Ala Pro Ala Val Ala Thr * 115 120

<210> 1055

<211> 122

<212> PRT

<213> Homo sapiens

<400> 1055

Met Leu Thr Cys Leu Phe Ser Phe Gln Gly Cys Trp Arg Ala Arg Gly 1.0 Trp Gln Arg Leu Cys Glu Gly Arg Arg Gly Trp Pro Gly Val Gly Gln 2.5 Arg Thr Leu Lys Val Ser Glu Pro Ala Pro Leu Arg Val Gly Arg Ala 40 Leu Pro Gln Ala Leu Leu Gly Ala Arg Pro His Cys Val Phe Pro Gly 55 60 Gly Glu Val Leu Gly Val Glu Ala Ala Phe Gly Ser Ser Phe Ile Leu 70 75 Ser Thr Phe Phe Leu His Gln Pro Leu Phe Phe Pro Gly Pro Lys Leu 85 90 Arg Ala Thr Gln Tyr Leu Ile Ser Ser Asp Pro Thr His Leu Pro Ala 105 Gly Arg Gly Pro Asn Ser Val Ser Met 115 120 121

<210> 1056

<211> 51

<212> PRT

<213> Homo sapiens

<400> 1056

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<210> 1057 <211> 260 <212> PRT <213> Homo sapiens

<400> 1057 Met Glu Ala Pro Ala Gln Leu Leu Phe Leu Leu Leu Trp Leu Pro 10 Asp Thr Thr Gly Glu Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser 25 Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser 40 Val Gly Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro 55 Arg Pro Leu Ile Tyr Asp Ala Ser Asn Arg Ala Thr Gly Ile Pro Ala 70 75 Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser 85 90 Ser Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln His Arg Asp 105 Asn Trp Pro Pro Gly Ala Thr Phe Gly Gly Gly Thr Lys Val Glu Ile 125 120 Lys His Thr Thr Gly Glu Ile Val Leu Thr Gln Ala Pro Gly Thr Leu 135 140 Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln 155 150 Thr Ile Gly Ser Thr Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys 170 175 Ala Pro Lys Leu Leu Ile Tyr Trp Phe Ile Gln Phe Ala Lys Arg Gly 185 Pro Ile Lys Val Gln Cys His Arg Val Arg Gly Gln Thr Ser Leu Ser 200 Pro Ser Ala Asp Trp Ser Leu Lys Ile Leu Gln Cys Ile Ser Val Thr 215 220 Asn Met Gly Ala His Pro Thr Leu Leu Ala Glu Gly Pro Arg Trp Arg 230 235 Ser Asn Glu Leu Trp Leu His His Leu Ser Ser Ser Arg His Leu 245 Met Ser Ser 259

<210> 1058 <211> 52 <212> PRT <213> Homo sapiens

Trp Arg Pro Cys Leu Pro Arg Leu Arg Met Arg Val Leu Val Leu Leu 35 40 45

Ile Trp Ser * 50 51

<210> 1059 <211> 97 <212> PRT <213> Homo sapiens

<400> 1059 Met Gly Arg Gly Ser Glu Leu Pro Val Cys Leu Ala Phe Leu Val Cys 5 10 Leu Met Ala Ala Leu Gly Cys Cys Glu Val Leu Ser Thr Val His Pro 20 25 Glu Glu Thr Val Leu Arg Ala Pro Pro Thr Asn Phe Gln Arg Cys Gln 40 Leu Gln Gln Gly Ser Ala Leu Val Arg Glu Thr Ala Trp Gly Val Gly 55 Arg Gly Arg Pro Ser Glu Arg Trp His Gly Glu Leu Ala Gly Gly 70 75 Ser Arg Arg Asp Gly Met Glu Gly Leu Gly Pro Val Leu Leu Gly Ala 90

<210> 1060 <211> 99 <212> PRT <213> Homo sapiens

<400> 1060 Met Asn Lys His Phe Leu Phe Leu Phe Leu Leu Tyr Cys Leu Ile Ala 10 Ala Val Thr Ser Leu Gln Cys Ile Thr Cys His Leu Arg Thr Arg Thr 20 25 Asp Arg Cys Arg Arg Gly Phe Gly Val Cys Thr Ala Gln Lys Gly Glu 35 40 Ala Cys Met Leu Leu Arg Ile Tyr Gln Arg Asn Thr Leu Gln Ile Ser 55 Tyr Met Val Cys Gln Lys Phe Cys Arg Asp Met Thr Phe Asp Leu Arg 70 Asn Arg Thr Tyr Val His Thr Cys Cys Asn Tyr Asn Tyr Cys Asn Phe 90 Lys Leu * 98

<210> 1061 <211> 64 <212> PRT <213> Homo sapiens

<210> 1062 <211> 149 <212> PRT <213> Homo sapiens

<400> 1062 Met Tyr Leu Ser Asn Thr Thr Val Thr Ile Leu Ala Asn Leu Val Pro 10 15 1 5 Phe Thr Leu Thr Leu Ile Ser Phe Leu Leu Ile Cys Ser Leu Cys 20 25 Lys His Leu Lys Lys Met Gln Leu His Gly Lys Gly Ser Gln Asp Pro 40 Ser Met Lys Val His Ile Lys Ala Leu Gln Thr Val Thr Ser Phe Leu 55 Leu Leu Cys Ala Ile Tyr Phe Leu Ser Met Ile Ile Ser Val Cys Asn 70 75 80 Phe Gly Arg Leu Glu Lys Gln Pro Val Phe Met Phe Cys Gln Ala Ile 90 Ile Phe Ser Tyr Pro Ser Thr His Pro Phe Ile Leu Ile Leu Gly Asn 105 Lys Lys Leu Lys Gln Ile Phe Leu Ser Val Leu Arg His Val Arg Tyr 115 120 Trp Val Lys Asp Arg Ser Leu Arg Leu His Arg Phe Thr Arg Gly Ala Leu Cys Val Phe *

<210> 1063 <211> 63 <212> PRT <213> Homo sapiens

148

145

<210> 1064 <211> 92 <212> PRT <213> Homo sapiens

<210> 1065 <211> 67 <212> PRT <213> Homo sapiens

> <210> 1066 <211> 78 <212> PRT <213> Homo sapiens

50 55 60
Leu Ala Gly Trp Asp Leu Thr Gly Ala Pro Gly Ser Leu Gly 65 70 75 78

<210> 1067 <211> 55

<212> PRT

<213> Homo sapiens

<400> 1067

Met Tyr Phe Gly Ala Tyr Ala Phe Thr Val Ala Pro Arg Leu Ala Ile 1 5 5 10 10 15 Leu Gln Val Val Asn Val Ile Ser Tyr Lys Asp Ile Arg His Phe Tyr 20 25 30 Leu Arg His Trp Arg Asn Glu Arg Asn Cys Ile Cys His Val Asp Gly 35 40 45 Ala Leu Ile Lys Glu Gln *

<210> 1068

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1068

<210> 1069

<211> 64

<212> PRT

<213> Homo sapiens

<400> 1069

<210> 1070

<211> 73 <212> PRT <213> Homo sapiens

<210> 1071 <211> 152 <212> PRT <213> Homo sapiens

<400> 1071 Met Phe Trp Thr Met Ile Ile Leu Gln Val Leu Ile Pro Ile Ser 5 10 Leu Tyr Val Ser Ile Glu Ile Val Lys Leu Gly Gln Ile Tyr Phe Ile 25 Gln Ser Asp Val Asp Phe Tyr Asn Glu Lys Met Asp Ser Ile Val Gln 40 Cys Arg Ala Leu Asn Ile Ala Glu Asp Leu Gly Gln Ile Gln Tyr Leu Phe Ser Asp Lys Thr Gly Thr Leu Thr Glu Asn Lys Met Val Phe Arg Arg Trp Ser Gly Gly Arg Phe Asp Tyr Cys Pro Gly Glu Lys Ala Arg 90 Arg Val Glu Ser Phe Gln Glu Ala Phe Glu Glu His Phe Leu 105 Thr Thr Gly Arg Gly Phe Leu Thr His Met Ala Asn Pro Arg Ala Pro 120 125 Pro Leu Ala Asp Thr Phe Lys Met Gly Ala Ser Gly Arg Leu Ser Pro 135 Pro Ser Leu Thr Ala Arg Gly Ala 150 152

<210> 1072 <211> 113 <212> PRT <213> Homo sapiens

<210> 1073

<211> 52

<212> PRT

<213> Homo sapiens

<400> 1073

 Met Thr Leu Cys
 Cys
 Pro Trp Ala Thr Met His Pro Ser Thr Val Leu

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 Arg Met Val Trp Ser Leu Arg Ser Arg Ala Arg Arg Trp Gly Ser Val
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 Arg Thr Gly Leu Ser Trp Ser Ser Ser Ser Asp Ser Arg Ile Thr Ser
 45

 Leu Ser Leu *
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 51

<210> 1074

<211> 78

<212> PRT

<213> Homo sapiens

<400> 1074

 Met
 Phe
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 Arg
 Leu
 Tyr
 Ala
 Val
 Cys
 Met
 Leu
 Tyr
 Ala
 Val
 Cys
 Met
 Leu
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 Val
 Asp
 Lys
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<210> 1075

<211> 253

<212> PRT

<213> Homo sapiens

<400> 1075 Met Ser Ser Pro Gly Leu Leu Phe Ser Ser Leu Ser His Leu Leu 10 5 Leu Asn Ser Ser Thr Leu Ala Leu Leu Thr His Arg Leu Ser Gln Met 20 25 Thr Cys Leu Gln Ser Leu Arg Leu Asn Arg Asn Ser Ile Gly Asp Val 40 Gly Cys Cys His Leu Ser Glu Ala Leu Arg Ala Ala Thr Ser Leu Glu Glu Leu Asp Leu Ser His Asn Gln Ile Gly Asp Ala Gly Asp Gln His 75 70 Leu Ala Thr Ile Leu Pro Gly Leu Pro Glu Leu Arg Lys Ile Asp Leu 85 90 Ser Gly Asn Ser Ile Ser Ser Ala Gly Gly Val Gln Leu Ala Glu Ser 105 Leu Val Leu Cys Arg Arg Leu Glu Glu Leu Met Leu Gly Cys Asn Ala . 125 120 Leu Gly Asp Pro Thr Ala Leu Gly Leu Ala Gln Glu Leu Pro Gln His 135 140 Leu Arq Val Leu His Leu Pro Phe Ser His Leu Gly Pro Asp Gly Ala 150 155 Leu Ser Leu Ala Gln Asp Leu Asp Gly Ser Pro His Leu Glu Glu Ile 170 165 Ser Leu Ala Glu Asn Asn Leu Ala Gly Gly Val Leu Arg Phe Cys Met 185 Glu Leu Pro Leu Leu Arg Gln Ile Glu Leu Ser Trp Asn Leu Leu Gly 200 205 Asp Glu Ala Ala Glu Leu Ala Gln Val Leu Pro Gln Met Gly Arg 215 220 Leu Lys Arg Val Glu Tyr Glu Gly Pro Gly Glu Glu Trp Asp Gly Leu 230 235 Lys Gly Asp Leu His Pro Gly Asn Thr Lys Arg Pro Leu 245 250

<210> 1076 <211> 64 <212> PRT <213> Homo sapiens

<210> 1077 <211> 147 <212> PRT <213> Homo sapiens

<400> 1077 Met Met Lys Ser Leu Arg Val Leu Leu Val Ile Leu Trp Leu Gln Leu 10 Ser Trp Val Trp Ser Gln Gln Lys Glu Val Glu Gln Asn Ser Gly Pro 20 25 Leu Ser Val Pro Glu Gly Ala Ile Ala Ser Leu Asn Cys Thr Tyr Ser 40 Asp Arg Gly Ser Gln Ser Phe Phe Trp Tyr Arg Gln Tyr Ser Gly Lys 55 Ser Pro Glu Leu Ile Met Ser Ile Tyr Ser Asn Gly Asp Lys Glu Asp 70 75 Gly Arg Phe Thr Ala Gln Leu Asn Lys Ala Ser Gln Tyr Val Ser Leu 90 Leu Ile Arg Asp Ser Gln Pro Ser Asp Ser Ala Thr Tyr Leu Cys Ala 100 105 Asp Tyr Ser Gly Asn Thr Pro Leu Val Phe Gly Lys Gly Thr Arg Leu 120 125 Ser Val Ile Ala Asn Ile Gln Asn Pro Asp Pro Ala Leu Tyr Gln Leu Arg Asp Ser 145 147

<210> 1078

<211> 55

<212> PRT

<213> Homo sapiens

<400> 1078

 Met
 Phe
 Gln
 Gly
 Ser
 Asn
 Ile
 Leu
 Phe
 Leu
 Pro
 Ser
 Pro
 Gly
 Ile

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 Thr
 Ser
 Ile
 Asn
 Asp
 Arg
 Thr
 Tyr
 Phe
 Leu
 Phe
 Val
 Met
 Arg
 Ser
 Asn

 20
 20
 25
 25
 30
 30
 Asn
 Lys

 Trp
 Leu
 Phe
 Leu
 Lys
 Asn
 Lys
 Asn
 Lys

 Ser
 Leu
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 Leu
 Lys
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 45

 Ser
 Leu
 Lys
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 50
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<210> 1079

<211> 97

<212> PRT

<213> Homo sapiens

<400> 1079

 Met Ile Pro Ala Phe Gly Ile Phe Arg Leu Leu Ile Ile Ile Leu Ile

 1
 5
 10
 15

 Ile Val Leu Asp Met Gly Phe Ala Leu Tyr Arg Arg Phe Phe Val Pro
 20
 25
 30

 Glu Asp Gly Ser Pro Val Ser Phe Ala Ala His Ile Ala Gly Gly Phe
 45

 Ala Gly Met Ser Ile Gly Tyr Thr Val Phe Ser Cys Phe Asp Lys Ala
 50

Leu Met Lys Asp Pro Arg Phe Trp Ile Ala Ile Ala Ala Tyr Leu Ala 65 70 75 80

Cys Val Leu Phe Ala Val Phe Phe Asn Ile Phe Leu Ser Pro Ala Asn 85 90 95 96

<210> 1080 <211> 134 <212> PRT <213> Homo sapiens

. <400> 1080 Met Leu Ser Ile Leu Leu Ala Thr Leu Thr Leu Ser Leu Lys Glu Lys 10 5 Arg Gly Glu Arg Ser Ile His Gln Pro Glu Pro Ser Glu Lys Ser Val 25 Cys Leu Pro Val Ser Gly Ala Asp Pro Phe Arg Gly Ser Arg Gly Arg 40 Gly Lys Glu Ile Arg Arg Glu Lys Asp Ile Gly Leu Leu Glu His Val Gly Gln Glu Val Pro Arg Arg Ile Cys Glu Gln Leu Pro Asp Ser Lys 75 70 Ala Leu Ala Arg Pro Gln Asp Gly Pro Cys Leu Leu Asp Ile Arg Lys 90 Pro Lys Gly Gln Asn Lys Asn Thr Cys Leu Val Gly Glu Gly Ser Leu 105 Arg Gly His Gln Val Gly Gln Ile Pro Leu Val Thr His Leu Trp Arg 120 Leu Pro Gln Lys Cys * 130 133

<210> 1081 <211> 185 <212> PRT <213> Homo sapiens

<400> 1081 Met Lys Ile Leu Val Ala Phe Leu Val Val Leu Thr Ile Phe Gly Ile 10 Gln Ser His Gly Tyr Glu Val Phe Asn Ile Ile Ser Pro Ser Asn Asn 2.0 25 Gly Gly Asn Val Gln Glu Thr Val Thr Ile Asp Asn Glu Lys Asn Thr 40 Ala Ile Ile Asn Ile His Ala Gly Ser Cys Ser Ser Thr Thr Ile Phe 55 Asp Tyr Lys His Gly Tyr Ile Ala Ser Arg Val Leu Ser Arg Arg Ala 70 75 Cys Phe Ile Leu Lys Met Asp His Gln Asn Ile Pro Pro Leu Asn Asn 90 85 Leu Gln Trp Tyr Ile Tyr Glu Lys Gln Ala Leu Asp Asn Met Phe Ser 105 Ser Lys Tyr Thr Trp Val Lys Tyr Asn Pro Leu Glu Ser Leu Ile Lys

115 120 Asp Val Asp Trp Phe Leu Leu Gly Ser Pro Ile Glu Lys Leu Cys Lys 135 140 His Ile Pro Leu Tyr Lys Gly Glu Val Val Glu Asn Thr His Asn Val 150 155 Gly Ala Gly Gly Cys Ala Lys Ala Gly Leu Leu Gly Ile Leu Gly Ile 165 170 Ser Ile Cys Ala Asp Ile His Val * 180 184

<210> 1082 <211> 285 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(285)

<223> Xaa = any amino acid or nothing

<400> 1082 Met Val Ile Ala Leu Ile Ile Phe Leu Arg Ser Pro Ala Met Ala Gly 1 5 10 Gly Leu Phe Ala Ile Glu Arg Glu Phe Phe Blu Leu Gly Leu Tyr 20 25 Asp Pro Gly Leu Gln Ile Trp Gly Gly Glu Asn Phe Glu Ile Ser Tyr 40 Lys Ile Trp Gln Cys Gly Gly Lys Leu Leu Phe Xaa Pro Cys Ser Arg 55 60 Val Gly His Ile Tyr Arg Leu Glu Gly Trp Gln Gly Asn Pro Pro Pro 70 75 Ile Tyr Val Gly Ser Ser Pro Thr Leu Lys Asn Tyr Val Arg Val Val 85 90 Glu Val Trp Trp Asp Glu Tyr Lys Asp Tyr Phe Tyr Ala Ser Arg Pro 100 105 Glu Ser Gln Ala Leu Pro Tyr Gly Asp Ile Ser Glu Leu Lys Lys Phe 120 Arg Glu Asp His Asn Cys Lys Ser Phe Lys Trp Phe Met Glu Glu Ile 130 135 140 Ala Tyr Asp Ile Thr Ser His Tyr Pro Leu Pro Pro Lys Asn Val Asp 150 155 Trp Gly Glu Ile Arg Gly Phe Glu Thr Ala Tyr Cys Ile Asp Ser Met 170 Gly Lys Thr Asn Gly Gly Phe Val Glu Leu Gly Pro Cys His Arg Met 185 Gly Gly Asn Gln Leu Phe Arg Ile Asn Glu Ala Asn Gln Leu Met Gln 200 Tyr Asp Gln Cys Leu Thr Lys Gly Ala Asp Gly Ser Lys Val Met Ile 215 220 Thr His Cys Asn Leu Asn Glu Phe Lys Glu Trp Gln Tyr Phe Lys Asn 230 235 Leu His Arg Phe Thr His Ile Pro Ser Gly Lys Cys Leu Asp Arg Ser 250 245 Glu Val Leu His Gln Val Phe Ile Ser Asn Cys Asp Ser Ser Lys Thr 260 265 Thr Gln Lys Trp Glu Met Asn Asn Ile His Ser Val 275

280

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<210> 1083
<211> 73
<212> PRT
<213> Homo sapiens
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<210> 1084 <211> 56 <212> PRT <213> Homo sapiens

<210> 1085 <211> 68 <212> PRT <213> Homo sapiens

PCT/US01/02687 WO 01/54477

<210> 1086 <211> 62 <212> PRT <213> Homo sapiens

<400> 1086 Met Cys Pro Phe Met Pro Pro Pro Gly Leu Leu Arg Leu Phe Gln Ile 5 10 Val Phe Trp Val Glu His Pro Gly Ser Val Asn Pro Phe Glu Arg Ser Thr Ile Ile Gly Arg Ser Ala Lys Leu Lys Lys Asp Leu Lys Ser His Trp Glu Pro Gly Gln Gln Ala Leu Gln Gln Gly Leu Leu

<210> 1087 <211> 294 <212> PRT <213> Homo sapiens

<400> 1087 Met Pro Tyr Val Thr Glu Ala Thr Arg Val Gln Leu Val Leu Pro Leu 1 5 10 Leu Val Ala Glu Ala Ala Ala Pro Ala Phe Leu Glu Ala Phe Ala 25 Ala Asn Val Leu Glu Pro Arg Glu His Ala Leu Leu Thr Leu Leu Leu 40 Val Tyr Gly Pro Arg Glu Gly Gly Arg Gly Ala Pro Asp Pro Phe Leu 55 Gly Val Lys Ala Ala Ala Glu Leu Glu Arg Arg Tyr Pro Gly Thr 70 Arg Leu Ala Trp Leu Ala Val Arg Ala Glu Ala Pro Ser Gln Val Arg 85 90 Leu Met Asp Val Val Ser Lys Lys His Pro Val Asp Thr Leu Phe Phe 100 105 Leu Thr Thr Val Trp Thr Arg Pro Gly Pro Glu Val Leu Asn Arg Cys 120 Arg Met Asn Ala Ile Ser Gly Trp Gln Ala Phe Phe Pro Val His Phe 135 Gln Glu Phe Asn Pro Ala Leu Ser Pro Gln Arg Ser Pro Pro Gly Pro 155 Pro Gly Ala Gly Pro Asp Pro Pro Ser Pro Pro Gly Ala Asp Pro Ser 165 170 Arg Gly Ala Pro Ile Gly Gly Arg Phe Asp Arg Gln Ala Ser Ala Glu 185 Gly Cys Phe Tyr Asn Ala Asp Tyr Leu Ala Ala Arg Ala Arg Leu Ala 200 Gly Glu Leu Ala Gly Gln Glu Glu Glu Ala Leu Glu Gly Leu Glu 215 220 Val Met Asp Val Phe Leu Arg Phe Ser Gly Leu His Leu Phe Arg Ala 230 235 Val Glu Pro Gly Leu Val Gln Lys Phe Ser Leu Arg Asp Cys Ser Pro 245

<210> 1088 <211> 477 <212> PRT <213> Homo sapiens

<400> 1088 Met Gln Trp Lys Val Thr Leu Thr Ser Arg Trp Gly Leu Leu Arg His 10 Cys Gln Val Leu Ala Gly Leu Leu His Leu Gly Asn Ile Gln Phe Ala 20 25 Ala Ser Glu Asp Glu Ala Gln Pro Cys Gln Pro Met Asp Asp Ala Lys 40 Tyr Ser Val Arg Thr Ala Ala Ser Leu Leu Gly Leu Pro Glu Asp Val 55 Leu Leu Glu Met Val Gln Ile Lys Thr Ile Arg Ala Gly Arg Gln Gln 70 75 Gln Val Phe Arg Lys Pro Cys Ala Arg Ala Glu Cys Asp Thr Arg Arg 90 Asp Cys Leu Ala Lys Leu Ile Tyr Ala Arg Leu Phe Asp Trp Leu Val 105 Ser Val Ile Asn Ser Ser Ile Cys Ala Asp Thr Asp Ser Trp Thr Thr 120 Phe Ile Gly Leu Leu Asp Val Tyr Gly Phe Glu Ser Phe Pro Asp Asn 135 Ser Leu Glu Gln Leu Cys Ile Asn Tyr Ala Asn Glu Lys Leu Gln Gln 150 155 His Phe Val Ala His Tyr Leu Arg Ala Gln Glu Glu Tyr Ala Val 165 170 Glu Gly Leu Glu Trp Ser Phe Ile Asn Tyr Gln Asp Asn Gln Pro Cys 185 Leu Asp Leu Ile Glu Gly Ser Pro Ile Ser Ile Cys Ser Leu Ile Asn 200 205 Glu Glu Cys Arg Leu Asn Arg Pro Ser Ser Ala Ala Gln Leu Gln Thr 215 220 Arg Ile Glu Thr Ala Leu Ala Gly Ser Pro Cys Leu Gly His Asn Lys 235 230 Leu Ser Arg Glu Pro Ser Phe Ile Val Val His Tyr Ala Gly Pro Val 245 250 Arg Tyr His Thr Ala Gly Leu Val Glu Lys Asn Lys Asp Pro Ile Pro 265 Pro Glu Leu Thr Arg Leu Leu Gln Gln Ser Gln Asp Pro Leu Leu Met 280 Gly Leu Phe Pro Thr Asn Pro Lys Glu Lys Thr Gln Glu Glu Pro Pro 295 Gly Gln Ser Arg Ala Pro Val Leu Thr Val Val Ser Lys Phe Lys Ala 310 315 Ser Leu Glu Gln Leu Leu Gln Val Leu His Ser Thr Thr Pro His Tyr 325 330 Ile Arg Cys Ile Met Pro Asn Ser Gln Gly Gln Ala Gln Thr Phe Leu

345 Gln Glu Glu Val Leu Ser Gln Leu Glu Ala Cys Gly Leu Val Glu Thr 360 365 Ile His Ile Ser Ala Ala Gly Phe Pro Ile Arg Val Ser His Arg Asn 375 380 Phe Val Glu Arg Tyr Lys Leu Leu Arg Arg Leu His Pro Cys Thr Ser 390 395 Ser Gly Pro Asp Ser Pro Tyr Pro Ala Lys Gly Leu Pro Glu Trp Cys 405 410 Pro His Ser Glu Glu Ala Thr Leu Glu Pro Leu Ile Gln Asp Ile Leu 420 425 His Thr Leu Pro Val Leu Thr Gln Ala Ala Ile Thr Gly Asp Ser 440 Ala Glu Ala Met Pro Ala Pro Met His Cys Gly Arg Thr Lys Val Phe 455 Met Thr Asp Ser Met Leu Glu Leu Leu Glu Cys Gly Ala 475

<210> 1089

<211> 66

<212> PRT

<213> Homo sapiens

<400> 1089

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 Ala
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 Thr
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 Met

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<210> 1090

<211> 185

<212> PRT

<213> Homo sapiens

<400> 1090

Met Leu Trp Leu Leu Phe Phe Leu Val Thr Ala Ile His Ala Glu Leu 10 Cys Gln Pro Gly Ala Glu Asn Ala Phe Lys Val Arg Leu Ser Ile Arg 2.0 25 Thr Ala Leu Gly Asp Lys Ala Tyr Ala Trp Asp Thr Asn Glu Glu Tyr 40 Leu Phe Lys Ala Met Val Ala Phe Ser Met Arg Lys Val Pro Asn Arg 55 . 60 Glu Ala Thr Glu Ile Ser His Val Leu Leu Cys Asn Val Thr Gln Arg 70 75 Val Ser Phe Trp Phe Val Val Thr Asp Pro Ser Lys Asn His Thr Leu 85 90

Pro Ala Val Glu Val Gln Ser Ala Ile Arg Met Asn Lys Asn Arg Ile 100 105 Asn Asn Ala Phe Phe Leu Asn Asp Gln Thr Leu Glu Phe Leu Lys Ile 120 125 Pro Ser Thr Leu Ala Pro Pro Met Asp Pro Ser Val Pro Ile Trp Ile 140 135 Ile Ile Phe Gly Val Ile Phe Cys Ile Ile Ile Val Ala Ile Ala Leu 150 155 Leu Ile Leu Ser Gly Ile Trp Gln Arg Arg Lys Asn Lys Glu Pro 165 170 Ser Glu Val Asp Asp Ala Glu Glu * 180

<210> 1091 <211> 47 <212> PRT <213> Homo sapiens

<210> 1092 <211> 46 <212> PRT <213> Homo sapiens

<210> 1093 <211> 64 <212> PRT <213> Homo sapiens

35 40 45
Ser Leu Pro Gly Ala Pro Ala Thr Ser Ala Ser Pro Ser Val Leu *
50 55 60 63

<210> 1094 <211> 85 <212> PRT <213> Homo sapiens

<210> 1095 <211> 89 <212> PRT <213> Homo sapiens

<210> 1096 <211> 158 <212> PRT <213> Homo sapiens

Lys Phe Leu Lys Lys Ala Asp Thr Arg Asp Ser Arg Gln Ala Cys Leu Ala Ala Ser Leu Ala Leu Ala Leu Asn Gly Val Phe Thr Asn Thr Ile Lys Leu Ile Val Gly Arg Pro Arg Pro Asp Phe Phe Tyr Arg Cys Phe 55 Pro Asp Gly Leu Ala His Ser Asp Leu Met Cys Thr Gly Asp Lys Asp Val Val Asn Glu Gly Arg Lys Ser Phe Pro Ser Gly His Ser Ser Phe 85 90 Ala Phe Ala Gly Leu Ala Phe Ala Ser Phe Tyr Leu Ala Gly Lys Leu 100 105 110 His Cys Phe Thr Pro Gln Gly Arg Gly Lys Ser Trp Arg Phe Cys Ala 120 125 Phe Leu Ser Pro Leu Leu Phe Ala Ala Val Ile Ala Leu Ser Arg Thr 135 Cys Asp Tyr Lys His His Trp Gln Gly Pro Phe Lys Trp * 150

<210> 1097 <211> 88 <212> PRT

<213> Homo sapiens

Glu Pro Gln Leu Gly Gly Gly * 85 87

<210> 1098

<211> 58

<212> PRT

<213> Homo sapiens

<400> 1098

 Met
 Met
 Ser
 Gly
 Trp
 Leu
 Leu
 Arg
 Ala
 Ala
 Ile
 Cys
 Arg
 Gly
 Leu
 Leu

 Ser
 Ser
 Glu
 Ser
 Leu
 Thr
 Phe
 Thr
 Ser
 Ala
 Pro
 His
 Ser
 Ile
 Ser
 Ile

 Ser
 Ser
 Arg
 Asp
 Gly
 Asn
 Leu
 Gln
 Thr
 Gly
 Tyr
 Arg
 Pro
 Thr

 His
 Val
 Val
 Phe
 Leu
 Ser
 Thr
 Ala
 Arg
 *

 50
 55
 57
 57

<210> 1099 <211> 72 <212> PRT <213> Homo sapiens

<210> 1100 <211> 47 <212> PRT <213> Homo sapiens

<210> 1101 <211> 130 <212> PRT <213> Homo sapiens

<400> 1101 Met Arg Pro Leu Lys Pro Gly Ala Pro Leu Pro Ala Leu Phe Leu Leu 5 10 Ala Leu Ala Leu Ser Pro His Gly Ala His Gly Arg Pro Arg Gly Arg 20 25 Arg Gly Ala Arg Val Thr Asp Lys Glu Pro Lys Pro Leu Leu Phe Leu 40 Pro Ala Ala Gly Ala Gly Arg Thr Pro Ser Gly Ser Arg Ser Ala Glu 55 Ile Phe Pro Arg Asp Ser Asn Leu Lys Asp Lys Phe Ile Lys His Phe 70 75 Thr Gly Pro Val Thr Phe Ser Pro Glu Cys Ser Lys His Phe His Arg 85 90 Leu Tyr Tyr Asn Thr Arg Glu Cys Ser Thr Pro Ala Tyr Tyr Lys Arg 100 105

Cys Ala Arg Leu Leu Thr Arg Leu Ala Val Ser Pro Leu Cys Ser Gln
115
120
125
Thr *

<210> 1102 <211> 170 <212> PRT <213> Homo sapiens

<400> 1102 Met Gln Phe Val Leu Leu Arg Thr Leu Ala Tyr Ile Pro Thr Pro Ile 5 10 Tyr Phe Gly Ala Val Ile Asp Thr Thr Cys Met Leu Trp Gln Glu 20 25 Cys Gly Val Gln Gly Ser Cys Trp Glu Tyr Asn Val Thr Ser Phe Arg 40 Phe Val Tyr Phe Gly Leu Ala Ala Val Leu Lys Tyr Val Gly Cys Ile 55 60 Phe Ile Leu Leu Ala Trp Tyr Ser Ile Lys Asp Thr Glu Asp Glu Gln 70 75 Pro Arg Leu Arg Gln Lys Lys Ile Cys Leu Ser Thr Leu Ser Asp Thr 90 Met Thr Gln Pro Asp Ser Ala Gly Val Val Ser Cys Pro Leu Phe Thr 105 Pro Asp Gly Glu Ile His Lys Lys Thr Gly Leu Arg Lys Arg Asp Pro 120 Gly Gly Thr Thr Glu Pro Thr Pro Gly Pro Leu Arg Lys Arg Pro Leu 135 140 Cys Thr Leu Glu Ala Pro Arg Leu Pro Asn Lys Ala Pro Phe Thr Leu 150 155 Glu Leu Ala Leu Leu Arg Val Arg Leu * 165 169

<210> 1103 <211> 62 <212> PRT <213> Homo sapiens

<210> 1104 <211> 83

<212> PRT <213> Homo sapiens

<210> 1105 <211> 124 <212> PRT <213> Homo sapiens

<400> 1105 Met Val Phe Thr Val Thr Leu Lys Leu Ala Leu Asp Thr His Tyr Trp 1 5 10 Thr Trp Ile Asn His Phe Val Ile Trp Gly Ser Leu Leu Phe Tyr Val 25 Val Phe Ser Leu Leu Trp Gly Gly Val Ile Trp Pro Phe Leu Asn Tyr 40 Gln Arg Met Tyr Tyr Val Phe Ile Gln Met Leu Ser Ser Gly Pro Ala 55 Trp Leu Ala Ile Val Leu Leu Val Thr Ile Ser Leu Leu Pro Asp Val 70 75 Leu Lys Lys Val Leu Cys Arg Gln Leu Trp Pro Thr Ala Thr Glu Arg 85 90 Val Gln Thr Lys Ser Gln Cys Leu Ser Val Glu Gln Ser Thr Ile Phe 105 Met Leu Ser Gln Thr Ser Ser Ser Leu Ser Phe *

120 123

<210> 1106 <211> 248 <212> PRT <213> Homo sapiens

Leu Glu Ser Ser Trp Pro Phe Trp Leu Thr Leu Ala Leu Ala Val Ile 55 Leu Gln Asn Met Ala Ala His Trp Val Phe Leu Glu Thr His Asp Gly His Pro Gln Leu Thr Asn Arg Arg Val Leu Tyr Ala Ala Thr Phe Leu Leu Phe Pro Leu Asn Val Leu Val Gly Ala Met Val Ala Thr Trp Arg 100 105 110 Val Leu Leu Ser Ala Leu Tyr Asn Ala Ile His Leu Gly Gln Met Asp 120 125 Leu Ser Leu Leu Pro Pro Arg Ala Ala Thr Leu Asp Pro Gly Tyr Tyr 135 140 Thr Tyr Arg Asn Phe Leu Lys Ile Glu Val Ser Gln Ser His Pro Ala 150 155 Met Thr Ala Phe Cys Ser Leu Leu Leu Gln Ala Gln Ser Leu Leu Pro 165 170 Arg Thr Met Ala Ala Pro Gln Asp Ser Leu Arg Pro Gly Glu Glu Asp 180 185 Glu Gly Met Gln Leu Gln Thr Lys Asp Ser Met Ala Lys Gly Ala 200 Arg Pro Gly Ala Ser Arg Gly Arg Ala Arg Trp Gly Leu Ala Tyr Thr 215 220 Leu Leu His Asn Pro Thr Leu Gln Val Phe Arg Lys Thr Ala Leu Leu 235 230 Gly Ala Asn Gly Ala Gln Pro 245

<210> 1107 <211> 121 <212> PRT <213> Homo sapiens

Ala Phe Val Val Pro Glu His Gln Leu Gly Thr Ala Tyr Gly Phe Met 35 40 45 Gln Ser Ile Gln Asn Leu Gly Leu Ala Ile Ile Ser Ile Ile Ala Gly

50 55 60 Met Ile Leu Asp Ser Arg Gly Tyr Leu Phe Leu Glu Val Phe Phe Ile

65 70 75 80
Ala Cys Val Ser Leu Ser Leu Leu Ser Val Val Leu Leu Tyr Leu Val

Asn Arg Ala Gln Gly Gly Asn Leu Asn Tyr Ser Ala Arg Gln Arg Glu
100 105 110

Glu Ile Lys Phe Ser His Thr Glu * 115 120

<210> 1108 <211> 53 <212> PRT <213> Homo sapiens

<210> 1109 <211> 259 <212> PRT <213> Homo sapiens

<400> 1109 Met His Val Val Ile Val Leu Lys Ala Leu Val Ala Val Gln Ile Leu 1 5 10 Leu Ser Ile Lys Glu Tyr Thr Leu Glu Arg Asn His Met His Val Ile 20 25 Ser Val Ile Lys Val Leu Val Lys Ala Gln Thr Ser Leu Asn Ile Arg 40 Glu Tyr Thr Leu Val Lys Ser Leu Ile Ile Ala Ile Val Val Arg Lys 55 60 Pro Ser Val Arg Val Leu Thr Leu Phe Phe Ile Arg Glu Phe Thr Leu 70 75 Glu Lys Asn Tyr Tyr Leu Cys Thr Gln Cys Ser Lys Ser Phe Ser Gln 90 85 Ile Ser Asp Leu Ile Lys His Gln Arg Ile His Thr Gly Glu Lys Pro 100 105 Tyr Lys Cys Ser Glu Cys Arg Lys Ala Phe Ser Gln Cys Ser Ala Leu 120 125 Thr Leu His Gln Arg Ile His Thr Gly Lys Lys Pro Asn Pro Cys Asp 135 140 Glu Cys Gly Lys Ser Phe Ser Arg Arg Ser Asp Leu Ile Asn His Gln 150 155 Lys Ile His Thr Gly Glu Lys Pro Tyr Lys Cys Asp Ala Cys Gly Lys 165 170 Ala Phe Ser Thr Cys Thr Asp Leu Ile Glu His Gln Lys Thr His Ala 185 Glu Glu Lys Pro Tyr Gln Cys Val Gln Cys Ser Arg Ser Cys Ser Gln 200 Leu Ser Glu Leu Thr Ile His Glu Glu Val His Cys Gly Glu Asp Ser 215 220 Gln Asn Val Met Asn Val Arg Lys Pro Leu Val Cys Thr Pro Thr Leu 230 235 Phe Ser Thr Arg Asp Thr Val Pro Glu Lys Asn Leu Met Asn Ala Val 250 245 Asp Tyr *

<210> 1110

<211> 47 <212> PRT <213> Homo sapiens

<400> 1110

 Met Thr Cys
 Ser Leu Leu Ser Leu Leu Asp Ala Val Cys
 Ser Ser Phe

 1
 5
 10
 15

 Val Gln Ala Phe Cys
 Ser Arg Asp Pro Glu Arg Trp Pro Ala Ile Ser
 30

 Pro His Ser Leu Ser Gly Ala Phe Tyr Phe Leu Asn Val Cys
 *

 35
 45
 46

<210> 1111 <211> 93 <212> PRT <213> Homo sapiens

<400> 1111

 Met
 Ser
 Leu
 Arg
 Ala
 Pro
 Ser
 Val
 Arg
 Ile
 Phe
 Val
 Tyr
 Leu
 Leu
 Phe

 Arg
 Leu
 5
 10
 15
 15

 Arg
 Leu
 His
 Thr
 Gln
 Arg
 Gly
 Leu
 Ala
 Gly
 Arg
 Arg
 Arg
 Gln
 Trp
 Gly
 Ser
 Trp
 Ala
 Bly
 Arg
 Arg
 Arg
 Gln
 Trp
 Gly
 Arg
 Arg
 Arg
 Gln
 Trp
 Gly
 Ser
 Arg
 Arg
 Arg
 Gln
 Leu
 Ser
 Arg
 His
 Leu
 Ser
 Trp
 Arg
 Phe
 Pro
 Gly
 Trp
 Ala
 Ser
 Arg
 Phe
 Pro
 Cys
 Trp
 Arg
 Phe
 Pro
 Cys
 Trp
 Arg
 Phe
 Pro
 Cys
 Trp
 Arg
 Phe
 Pro</

<210> 1112 <211> 71 <212> PRT <213> Homo sapiens

<210> 1113 <211> 47 <212> PRT <213> Homo sapiens

<210> 1114 <211> 55 <212> PRT <213> Homo sapiens

<210> 1115 <211> 83 <212> PRT <213> Homo sapiens

<210> 1116 <211> 145 <212> PRT <213> Homo sapiens

<400> 1116 Met Val Leu Leu Val Val Gly Asn Leu Val Asn Trp Ser Phe Ala Leu Phe Gly Leu Ile Tyr Arg Pro Arg Asp Phe Ala Ser Tyr Met Leu Gly Ile Phe Ile Cys Asn Leu Leu Leu Tyr Leu Ala Phe Tyr Ile Ile Met 40 Lys Leu Arg Ser Ser Glu Lys Val Leu Pro Val Pro Leu Phe Cys Ile Val Ala Thr Ala Val Met Trp Ala Ala Ala Leu Tyr Phe Phe Gln 70 75 Asn Leu Ser Ser Trp Glu Gly Thr Pro Ala Glu Ser Arg Glu Lys Asn 85 90 Arg Glu Cys Ile Leu Leu Asp Phe Phe Asp Asp His Asp Ile Trp His 100 105 Phe Leu Ser Ala Thr Ala Leu Phe Phe Ser Phe Leu Asp Leu Leu Thr 120 Leu Asp Asp Asp Leu Asp Val Val Arg Arg Asp Gln Ile Pro Val Phe

<210> 1117 <211> 139 <212> PRT <213> Homo sapiens

<400> 1117 Met Gly Asp Phe Ala Gly Val Asp Phe Val Phe Leu Val Val Cys Phe 10ء Ala Gln Arg Gln Gly Ala Ala Glu Ala Val Gly Ala Val Leu Ala Val 2.0 Leu Leu Cys Asp Thr Leu Leu Gly Val Thr Arg Leu Glu Gly Val Ile His Leu Pro Leu Tyr Phe Gly Leu Ser Gly Ile Glu Val Ile Gln Gln 55 Ala His Asn Arg Gly Ser Ser Arg Phe Gln Leu Leu Ile Arg Trp Arg 70 75 Glu Asp Glu Asp Arg Trp Cys Ser His Ser Ser Phe Asp Val His Leu 85 90 Gly Pro Leu Ala Glu Arg Pro His Val Ser Thr Gln Leu Leu Thr Val 105 Ile Ser Cys Lys Ile Phe Arg Leu Gln Ala Thr Asp Cys Glu Ser Lys 120 Phe Cys Pro Arg Ser Ser Ala Ala Glu Pro * 135 138

<210> 1118 <211> 194 <212> PRT <213> Homo sapiens

<400> 1118 Met Cys Leu Leu Phe Leu Leu Pro Arg Phe Pro Val Ser Trp Arg Ala 5 10 Gly Val Asp Gly Ala Ala Pro Ser Ser Gln Asp Leu Trp Arg Ile Arg 25 Ser Pro Cys Gly Asp Cys Glu Gly Phe Asp Val His Ile Met Asp Asp 40 Met Ile Lys Arg Ala Leu Asp Phe Arg Glu Ser Arg Glu Ala Glu Pro 55 His Pro Leu Trp Glu Tyr Pro Cys Arg Ser Leu Ser Glu Pro Trp Gln 70 75 Ile Leu Thr Phe Asp Phe Gln Gln Pro Val Pro Leu Gln Pro Leu Cys 85 90 Ala Glu Gly Thr Val Glu Leu Lys Arg Pro Gly Gln Ser His Ala Ala 100 105 Val Leu Trp Met Glu Tyr His Leu Thr Pro Glu Cys Thr Leu Ser Thr 120 115 Gly Leu Leu Glu Pro Ala Asp Pro Glu Gly Gly Cys Cys Trp Asn Pro 130 135 140 His Cys Lys Gln Ala Val Tyr Phe Phe Ser Pro Ala Pro Asp Pro Arg 150 155 Ala Leu Leu Gly Gly Pro Arg Thr Val Ser Tyr Ala Val Glu Phe His 170 175 165 Pro Asp Thr Gly Asp Ile Ile Met Glu Phe Arg His Ala Asp Thr Pro 185 Asp * 193

<210> 1119 <211> 118 <212> PRT <213> Homo sapiens

<400> 1119

Met Leu Val Leu Leu Pro Arg Ser Lys Ala Met Pro Leu Leu Ser Val Asn Val Thr Leu Ala Phe Phe Pro Arg Asn Lys Glu Ile Val Lys Tyr 25 Leu Leu Asn Gln Gly Ala Asp Val Thr Leu Arg Ala Lys Asn Gly Tyr 40 Thr Ala Phe Asp Leu Val Met Leu Leu Asn Asp Pro Asp Ile Phe Gly 55 Gly Glu Leu Ile Gly Phe Leu Ser Val Val Thr Glu Leu Val Arg Leu 75 70 Leu Ala Ser Val Phe Met Gln Val Asn Lys Asp Ile Gly Arg Arg Ser 85 90 His Gln Leu Pro Leu Pro His Ser Lys Val Pro Thr Ala Leu Glu His 100 105 Pro Ser Ala Ala Arg * 115 117

<210> 1120 <211> 842 <212> PRT

<213> Homo sapiens

<400> 1120 Met Leu Trp Gly Ser Gly Lys Cys Lys Ala Leu Thr Lys Phe Lys Phe 10 Val Phe Phe Leu Arg Leu Ser Arg Ala Gln Gly Gly Leu Phe Glu Thr Leu Cys Asp Gln Leu Leu Asp Ile Pro Gly Thr Ile Arg Lys Gln Thr 40 Phe Met Ala Met Leu Leu Lys Leu Arg Gln Arg Val Leu Phe Leu Leu 55 Asp Gly Tyr Asn Glu Phe Lys Pro Gln Asn Cys Pro Glu Ile Glu Ala 70 Leu Ile Lys Glu Asn His Arg Phe Lys Asn Met Val Ile Val Thr Thr 90 85 Thr Thr Glu Cys Leu Arg His Ile Arg Gln Phe Gly Ala Leu Thr Ala 100 105 Glu Val Gly Asp Met Thr Glu Asp Ser Ala Gln Ala Leu Ile Arg Glu 120 125 Val Leu Ile Lys Glu Leu Ala Glu Gly Leu Leu Leu Gln Ile Gln Lys 135 140 Ser Arg Cys Leu Arg Asn Leu Met Lys Thr Pro Leu Phe Val Val Ile 150 155 Thr Cys Ala Ile Gln Met Gly Glu Ser Glu Phe His Ser His Thr Gln 170 Thr Thr Leu Phe His Thr Phe Tyr Asp Leu Leu Ile Gln Lys Asn Lys 185 His Lys His Lys Gly Val Ala Ala Ser Asp Phe Ile Arg Ser Leu Asp 200 His Cys Gly Tyr Leu Ala Leu Glu Gly Val Phe Ser His Lys Phe Asp 215 Phe Glu Leu Gln Asp Val Ser Ser Val Asn Glu Asp Val Leu Leu Thr 230 235 Thr Gly Leu Leu Cys Lys Tyr Thr Ala Gln Arg Phe Lys Pro Lys Tyr 245 250 Lys Phe Phe His Lys Ser Phe Gln Glu Tyr Thr Ala Gly Arg Arg Leu 265 Ser Ser Leu Leu Thr Ser His Glu Pro Glu Glu Val Thr Lys Gly Asn 280 Gly Tyr Leu Gln Lys Met Val Ser Ile Ser Asp Ile Thr Ser Thr Tyr 300 295 Ser Ser Leu Leu Arg Tyr Thr Cys Gly Ser Ser Val Glu Ala Thr Arg 310 315 Ala Val Met Lys His Leu Ala Ala Val Tyr Gln His Gly Cys Leu Leu 325 330 Gly Leu Ser Ile Ala Lys Arg Pro Leu Trp Arg Gln Glu Ser Leu Gln 345 Ser Val Lys Asn Thr Thr Glu Gln Glu Ile Leu Lys Ala Ile Asn Ile Asn Ser Phe Val Glu Cys Gly Ile His Leu Tyr Gln Glu Ser Thr Ser 375 380 Lys Ser Ala Leu Ser Gln Glu Phe Glu Ala Phe Phe Gln Gly Lys Ser 390 395 Leu Tyr Ile Asn Ser Gly Asn Ile Pro Asp Tyr Leu Phe Asp Phe Phe 405 410 Glu His Leu Pro Asn Cys Ala Ser Ala Leu Asp Phe Ile Lys Leu Gly 425 Phe Tyr Gly Gly Ala Met Ala Ser Trp Glu Lys Ala Ala Glu Asp Thr

```
440
Gly Gly Ile His Met Glu Glu Ala Pro Glu Thr Tyr Ile Pro Ser Arg
          455
Ala Val Ser Leu Phe Phe Asn Trp Lys Gln Glu Phe Arg Thr Leu Glu
                470
                                  475
Val Thr Leu Arg Asp Phe Ser Lys Leu Asn Lys Gln Asp Ile Arg Tyr
             485
                              490
Leu Gly Lys Ile Phe Ser Ser Ala Thr Ser Leu Arg Leu Gln Ile Lys
          500 505 510
Arg Cys Ala Gly Val Ala Gly Ser Leu Ser Leu Val Leu Ser Thr Cys
            520
Lys Asn Ile Tyr Ser Leu Met Val Glu Ala Ser Pro Leu Thr Ile Glu
          535
                               540
Asp Glu Arg His Ile Thr Ser Val Thr Asn Leu Lys Thr Leu Ser Ile
     550 555 560
His Asp Leu Gln Asn Gln Arg Leu Pro Gly Gly Leu Thr Asp Ser Leu
          565 570 575
Gly Asn Leu Lys Asn Leu Thr Lys Leu Ile Met Asp Asn Ile Lys Met
         580 585 590
Asn Glu Glu Asp Ala Ile Lys Leu Ala Glu Gly Leu Lys Asn Leu Lys
                      600
Lys Met Cys Leu Phe His Leu Thr His Leu Ser Asp Ile Gly Glu Gly
                   615
Met Asp Tyr Ile Val Lys Ser Leu Ser Ser Glu Pro Cys Asp Leu Glu
                                 635
                630
Glu Ile Gln Leu Val Ser Cys Cys Leu Ser Ala Asn Ala Val Lys Ile
                              650
Leu Ala Gln Asn Leu His Asn Leu Val Lys Leu Ser Ile Leu Asp Leu
                           665
Ser Glu Asn Tyr Leu Glu Lys Asp Gly Asn Glu Ala Leu His Glu Leu
                       680
Ile Asp Arg Met Asn Val Leu Glu Gln Leu Thr Ala Leu Met Leu Pro
                    695
Trp Gly Cys Asp Val Gln Gly Ser Leu Ser Ser Leu Leu Lys His Leu
                                 715
Glu Glu Val Pro Gln Leu Val Lys Leu Gly Leu Lys Asn Trp Arg Leu
                              730
Thr Asp Thr Glu Ile Arg Ile Leu Gly Ala Phe Phe Gly Lys Asn Pro
                           745
Leu Lys Asn Phe Gln Gln Leu Asn Leu Ala Gly Asn Arg Val Ser Ser
                       760
Asp Gly Trp Leu Ala Phe Met Gly Val Phe Glu Asn Leu Lys Gln Leu
                    775
                                     780
Val Phe Phe Asp Phe Ser Thr Lys Glu Phe Leu Pro Asp Pro Ala Leu
                                 795
                790
Val Arg Lys Leu Ser Gln Val Leu Ser Lys Leu Thr Phe Leu Gln Glu
                              810
             805
Ala Arg Leu Val Gly Trp Gln Phe Asp Asp Asp Leu Ser Val Ile
       820
                          825
Thr Gly Ala Phe Lys Leu Val Thr Ala *
      835
                        840 841
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<210> 1121

<211> 90

<212> PRT

<213> Homo sapiens

<210> 1122 <211> 129 <212> PRT <213> Homo sapiens

<400> 1122 Met Phe Leu Leu Phe Trp Phe Ile Leu Ser Glu Gly Cys Pro Leu Leu 10 Glu Gln Leu Asn Ile Ser Trp Cys Asp Gln Val Thr Lys Asp Gly Ile 20 25 Gln Ala Leu Val Arg Gly Cys Gly Gly Leu Lys Ala Leu Phe Leu Lys 40 Gly Cys Thr Gln Leu Glu Asp Glu Ala Leu Lys Tyr Ile Gly Ala His 55 Cys Pro Glu Leu Val Thr Leu Asn Leu Gln Thr Cys Leu Gln Ile Thr 70 Asp Glu Gly Leu Ile Thr Ile Cys Arg Gly Cys His Lys Leu Gln Ser 85 90 Leu Cys Ala Ser Gly Cys Ser Asn Ile Thr Asp Ala Ile Leu Asn Ala 105 Leu Ser Gln Asn Cys Pro Arg Leu Ile Ile Leu Glu Val Ala Arg Cys 120 Ser 129

<210> 1123 <211> 243 <212> PRT <213> Homo sapiens

55 Ala Arg Val Leu Val Asp Gly Glu Glu His Val Gly Phe Leu Lys Thr 70 75 Asp Gly Ser Phe Val Val His Asp Ile Pro Ser Gly Ser Tyr Val Val 85 90 Glu Val Val Ser Pro Ala Tyr Arg Phe Asp Pro Val Arg Val Asp Ile 100 105 Thr Ser Lys Gly Lys Met Arg Ala Arg Tyr Val Asn Tyr Ile Lys Thr 115 120 Ser Glu Val Val Arg Leu Pro Tyr Pro Leu Gln Met Lys Ser Ser Gly 135 Pro Pro Ser Tyr Phe Ile Lys Arg Glu Ser Trp Gly Trp Thr Asp Phe 150 155 Leu Met Asn Pro Met Val Met Met Val Leu Pro Leu Leu Ile Phe 170 Val Leu Leu Pro Lys Val Val Asn Thr Ser Asp Pro Asp Met Arg Arg 185 Glu Met Glu Gln Ser Met Asn Met Leu Asn Ser Asn His Glu Leu Pro 200 Asp Val Ser Glu Phe Met Thr Arg Leu Phe Ser Ser Lys Ser Ser Gly 215 Lys Ser Ser Ser Gly Ser Ser Lys Thr Gly Lys Ser Gly Ala Gly Lys 230 235 Arg Arg * 242

<210> 1124

<211> 71

<212> PRT

<213> Homo sapiens

<400> 1124

 Met
 Leu
 Ser
 Tyr
 Ala
 His
 Ile
 Thr
 Leu
 Ala
 Val
 Leu
 Arg
 Ile
 Pro
 Ser

 Ala
 Thr
 Gly
 Cys
 Trp
 Arg
 Ala
 Phe
 Phe
 Thr
 Cys
 Ala
 Ser
 His
 Leu
 Thr
 Leu
 Thr
 Cys
 Ala
 Ser
 His
 Leu
 Thr
 Thr
 Thr
 Ala
 Leu
 Phe
 Met
 Tyr
 Val
 Arg
 Pro
 Arg
 Pro
 Arg
 Ser
 Asn
 Lys
 Leu
 Ile
 Ser
 Val
 Leu
 Tyr
 Arg
 Tyr
 Arg
 Ser
 Arg
 Ser
 Asn
 Lys
 Leu
 Ile
 Ser
 Val
 Leu
 Tyr
 Arg
 Ile
 Tyr
 Arg
 Tyr
 Arg
 Ser
 Arg
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 Arg
 Ser
 Arg
 Ser
 Arg
 Ser
 Arg
 Arg
 Arg
 Arg
 Arg

<210> 1125

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1125

Met Pro Thr Leu Gly Asp Ala Leu Ile Leu Tyr Leu His Leu Val Leu 1 5 10 15 Gly Val Ala Gly Val Leu Gln Pro Pro Gly Pro Arg Pro Ser Gln Ala 20 25 30

Leu Gly Pro Thr Gly Asp Arg Ala Pro Gly Lys Trp Asn Arg Ser * 35 40 45 47

<210> 1126 <211> 159 <212> PRT <213> Homo sapiens

<400> 1126 Met Phe Leu Ile Val Leu Pro Leu Glu Ser Met Ala His Gly Leu Phe 5 10 His Glu Leu Gly Asn Cys Leu Gly Gly Thr Ser Val Gly Tyr Ala Ile 20 25 Val Ile Pro Thr Asn Phe Cys Ser Pro Asp Gly Gln Pro Thr Leu Leu 40 45 Pro Pro Glu His Val Gln Glu Leu Asn Leu Arg Ser Thr Gly Met Leu 55 60 Asn Ala Ile Gln Arg Phe Phe Ala Tyr His Met Ile Glu Thr Tyr Gly 70 75 Cys Asp Tyr Ser Thr Ser Gly Leu Ser Phe Asp Thr Leu His Ser Lys 85 90 Leu Lys Ala Phe Leu Glu Leu Arg Thr Val Asp Gly Pro Arg His Asp 105 Thr Tyr Ile Leu Tyr Tyr Ser Gly His Thr His Gly Thr Gly Glu Trp 120 Ala Leu Ala Gly Gly Asp Thr Leu Arg Leu Asp Thr Leu Ile Glu Trp

140

155

135

150

Trp Arg Glu Lys Asn Gly Ser Phe Cys Ser Pro Pro Tyr Tyr Arg

<210> 1127 <211> 76 <212> PRT <213> Homo sapiens

<210> 1128 <211> 140 <212> PRT <213> Homo sapiens

<400> 1128 Met Gly Ala Gly Leu Ala Val Val Pro Leu Met Gly Leu Leu Glu Ser 5 10 Ile Ala Val Ala Lys Ala Phe Ala Ser Gln Asn Asn Tyr Arg Ile Asp 25 Ala Asn Gln Glu Leu Leu Ala Ile Gly Leu Thr Asn Met Leu Gly Ser Leu Val Ser Ser Tyr Pro Val Thr Gly Ser Phe Gly Arg Thr Ala Val 55 Asn Ala Gln Ser Gly Val Cys Thr Pro Ala Glu Gly Leu Val Thr Glu 70 75 Val Leu Val Leu Leu Ser Leu Asp Tyr Leu Thr Ser Leu Phe Tyr Tyr 85 90 Ile Pro Lys Ser Ala Leu Ala Ala Val Ile Ile Met Ala Val Ala Pro 100 105 Leu Phe Asp Thr Lys Ile Phe Arg Thr Leu Trp Arg Val Lys Arg Leu 115 120 Asp Leu Leu Ser Leu Ser Val Thr Phe Leu Leu Cys 130 135

<210> 1129 <211> 116 <212> PRT

<213> Homo sapiens

<400> 1129

Met Ala Glu Ala Phe Pro Phe Phe Ser Pro Phe Leu Gly Trp Leu Gly 5 10 Val Phe Leu Thr Gly Ser Asp Thr Ser Ser Asn Ala Leu Phe Ser Ser 25 Leu Gln Ala Thr Thr Ala His Gln Ile Gly Val Ser Asp Val Leu Leu 40 Val Ala Ala Asn Thr Ser Gly Gly Val Thr Gly Lys Met Ile Ser Pro Gln Ser Ile Ala Val Ala Cys Ala Ala Thr Gly Leu Val Gly Lys Glu Ser Asp Leu Phe Arg Phe Thr Leu Lys His Ser Leu Phe Phe Ala Thr 85 90 Ile Val Gly Leu Ile Thr Leu Ala Gln Ala Tyr Trp Phe Thr Gly Met 100 105 Leu Val His * 115

<210> 1130

<211> 81

<212> PRT

<213> Homo sapiens

<400> 1130

Met Asn Lys Leu Leu Val Ala Ala Thr Ala Ile Leu Phe Ser Leu Gly
1 5 10 15

Cys His Glu Lys Cys Lys Ile Phe Phe Leu Lys Ser Ile Ser Ser Pro
20 25 30

Gln Ser Leu Phe Leu Ala Asp Leu Cys Ala Ser Glu Pro Tyr Leu Leu
35 40 45

Phe Leu Asn Ala Val Leu Ser Ala Cys Asn Thr Ile Ser Phe Ile Ser
50 55 60

Val Pro Glu Ser Ser Gly Phe Ala Pro Ser Pro Pro Ala Ile Leu Leu
65 70 75 80

Leu
81

<210> 1131 <211> 46 <212> PRT <213> Homo sapiens

<210> 1132 <211> 46 <212> PRT <213> Homo sapiens

<210> 1133 <211> 87 <212> PRT <213> Homo sapiens

50 55 60

Glu Gln Ala Arg Glu Ser Leu Leu Ser Thr Phe Arg Ile Arg Pro Arg
65 70 75 80

Gly Arg Tyr Val Ser Tyr *
85 86

<210> 1134 <211> 57 <212> PRT <213> Homo sapiens

<210> 1135 <211> 57 <212> PRT <213> Homo sapiens

<210> 1136 <211> 105 <212> PRT <213> Homo sapiens

<210> 1137 <211> 52 <212> PRT <213> Homo sapiens

<210> 1138 <211> 187 <212> PRT <213> Homo sapiens

<400> 1138 Met Gln Pro Ile Val Ala Lys Ala Leu Val Val Leu Leu Glu Val His 5 10 Pro Leu Gln Asp Gln Ala Glu Ser Gly Arg Leu Gly His Val His Leu 20 25 Leu Cys Ala Pro Ala Ala Leu Gln His Ala Leu Arg Gly Ile Thr Leu His Asn Gly His His Gln Ala Asp His Leu Pro Asp Leu Met His His Glu Ala Leu Ala Leu His Pro Asp His Arg Lys Leu Gln Ala Leu Pro 70 75 His Lys Gly Phe Leu Ala Val His Leu Gln Asp Val Ala Ala Gly Thr 85 90 Gly Ile Leu Arg Pro Leu Leu Arg Gly Glu Ile Val Glu Val Val Arg 105 Ala Leu Val Ala Gly Gln Glu Pro Val Asp Leu Leu Gln Arg Leu Gly 120 125 Ala Gln Ala Val Gly Leu Ile Leu Asn Val Pro Val Leu Val Arg Lys 135 140 Gly Lys Arg Gly Gln Gln Val Ala Ile Gly Pro Gly Ile Thr Ser Val 150 155 160 Leu Gly Val Lys Pro Ala Arg Asp Pro Leu Gln Ser Gln Asn Pro Asn 165 170 Val Arg Gly Lys Val Ala Val Asp Leu Phe * 180 185 186

<210> 1139 <211> 109 <212> PRT <213> Homo sapiens

<400> 1139 Met Trp Gln Lys Ser Leu Leu Ile Leu Ser Phe Arg Val Ser Phe Pro 1 5 10 Leu Phe Leu Thr Tyr Asn Tyr Lys Leu Leu Ser Ile Arg Arg Thr Arg 25 20 Pro Leu Ser Ser Phe Phe Ser Lys Leu Leu Gln Ile Ala Val Asn Ser 40 Ile Asn Ser Leu Phe Ser Ala Gly Lys Val Ala Phe Ser Lys His Val 55 Cys Leu Leu Pro Gly Gly Leu Lys Ser Met Ile Tyr Cys Ser Ser Met 70 75 Cys Leu Lys Gln Leu Leu Arg Ser Phe Lys Gln Glu Ser Ser Lys Gly 90 Ser Val Leu Ile Met Val Leu Val Phe Leu Gln Ile * 105 108

<210> 1140 <211> 83 <212> PRT <213> Homo sapiens

<210> 1141 <211> 58 <212> PRT <213> Homo sapiens

Ser Ser Lys Phe Ser Trp Lys Ser Phe Ser Lys Leu Gln Phe Leu Leu 35 40 45

Leu Leu Lys Phe Arg Tyr Met Cys Ile *
50 55 57

<210> 1142 <211> 46 <212> PRT <213> Homo sapiens

<210> 1143 <211> 58 <212> PRT <213> Homo sapiens

<210> 1144 <211> 147 <212> PRT <213> Homo sapiens

 Arg
 His
 Ser
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<210> 1145 <211> 103 <212> PRT <213> Homo sapiens

<400> 1145 Met Ala Trp Ile Pro Leu Phe Leu Gly Val Leu Ala Tyr Cys Thr Gly 1 5 10 Ser Val Ala Ser Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ser 20 25 Pro Gly Lys Thr Ala Ser Ile Thr Cys Ser Gly Asp Lys Leu Gly Asp 40 Lys Tyr Ala Ser Trp Tyr Gln Gln Lys Ala Gly Gln Ser Pro Val Leu 55 Val Ile Tyr Glu Asp Ser Arg Arg Pro Ser Gly Ile His Lys Arg Phe 70 75 Tyr Gly Ser Asn Ser Gly Thr Thr Ala Thr Leu Thr Ile Ser Gly Thr 85 90 Gln Ala Met Asp Glu Gly * 100 102

<210> 1146 <211> 77 <212> PRT <213> Homo sapiens

<210> 1147 <211> 118 <212> PRT

<213> Homo sapiens

<400> 1147 Met Asn Pro Ser Ala Ser Leu Val Cys Leu Leu Phe Ala Phe Ser Ser 10 Cys Arg Ile Trp Ser Val Leu Cys Gln Leu Cys Val Pro Ser Pro Trp 25 Pro Ser Pro Leu Cys Leu Cys Pro Gln Thr Asp Val Ala Pro Ile Cys 3.5 40 Ala Val Gln Pro Ser Leu Phe Cys Leu Gly Ser Arg Glu Pro Leu Trp 55 60 Thr Val Leu Val Gly Ser Cys Pro Leu Arg Ala Phe Thr Asn Leu Ser 70 Val Arg Pro Pro Pro Gly His His Ser Ile His Leu Leu Thr Trp Leu 85 90 Ala Ser Ser Ser Ala Ala Ala Thr Thr Ala Ala Ser Thr Ala Ser Gly 100 105 Ala Pro His Ser Val * 115 117

<210> 1148 <211> 399 <212> PRT <213> Homo sapiens

<400> 1148 Met Trp Ala Ala Val Gly Gly Phe Leu Phe Ala Pro Arg Cys Phe Leu 10 Leu Pro Trp Pro Leu Arg Ala Pro Leu Ser Ser Leu Phe Val Leu Pro 20 25 Arg Leu Leu Trp Pro Ile Pro Tyr Pro Val Leu Ala Ser Val Cys 40 Pro Cys Val Pro Gly Gly Arg Phe Phe Gly Pro Leu Tyr Pro Arg Asp 55 Leu Arg Leu Arg Cys Val Pro Gly Glu Leu Thr Gly Ala Ala Pro 70 Arg Thr Leu Pro Gly Cys Asp Leu Asn Cys Leu Gly Leu Gly Arg Glu 85 90 Ala Ala Val Pro Arg Leu Leu Arg Leu Thr Arg Asp Pro Ala Arg Pro 105 Ser Cys Arg Thr Leu Gly Val His Ala Val Pro Arg Arg Ala Phe Gly 120 125 Phe Tyr Ala Val Pro Arg Arg Asp Pro Arg Phe Tyr Ala Val Pro Arg 140 135 Arg Val Pro Arg Leu Tyr Ala Val Pro His Pro Ala Leu Arg Val Tyr 150 155 Ala Val Pro Arg Arg Thr Phe Arg Val Tyr Ala Val Pro His Pro Ala 165 170 Leu Arg Val Tyr Ala Val Pro Arg Arg Ala Leu Gly Leu Tyr Val Val 180 185 Pro Gln Arg Ala Leu Arg Val Tyr Ala Val Pro Arg Arg Thr Phe Arg 200 Val Tyr Ala Val Pro His Pro Ala Leu Arg Leu Tyr Ala Val Ala Arg 215 Arg Ala Leu Arg Phe Tyr Val Val Pro Gln Arg Ala Leu Arg Val Tyr

230 235 Ala Val Pro Arg Leu Pro Gly Arg Ala Thr Phe Arg Asp Leu Arg Pro 245 250 Leu Leu Arg Leu Leu Pro Leu Gly Gly Arg Arg Val Leu Gly Leu 260 265 Pro Leu Ser Leu Pro Ala Gly Leu Ala Leu Arg Ala Ala Ser Arg Ala 280 Arg Pro Leu His Leu Leu Arg Ala Ala Cys Leu Leu Pro Ser Leu Gly 295 His Leu Gly Thr Leu Arg Gly Ser Leu Leu Gly Leu Ser Leu Ala Val 315 310 Arg Pro Pro Arg Ala Pro Arg Leu Gly Leu Arg Ala Pro Val Trp Pro 325 330 Ala Ala Ser Cys Leu Leu His Ser Gly Gly Ala Pro Arg Arg Leu Leu 340 345 350 Cys Ala Leu Ala Pro Leu Arg Pro Phe Cys Leu Pro Ala Arg Gly Ser 360 Trp Leu Ser Gly Ser Leu Ser Gln Arg Arg Gly Asp Leu Arg Arg Pro 375 380 Leu Gly Thr Arg Gly Asn Pro Leu Arg Leu Arg Gly Leu Gly His 390 395

<210> 1149 <211> 67 <212> PRT

<213> Homo sapiens

<400> 1149

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 Pro
 Ser
 Tyr
 Phe
 Lys
 Thr
 Cys
 Ser
 Leu
 Phe
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<210> 1150 <211> 70 <212> PRT

<213> Homo sapiens

<400> 1150

 Met
 Leu
 Val
 Ser
 Lys
 Leu
 Met
 Leu
 Gln
 Ile
 Val
 Met
 Ala
 Pro
 His

 Tyr
 Ile
 Met
 Pro
 Val
 Glu
 Met
 Lys
 Asn
 Gln
 Ser
 Leu
 Ile
 Pro
 Leu
 Leu
 Leu
 Leu
 Leu
 Leu
 Lys
 His
 Gly
 Glu
 Ser

 Leu
 Asp
 Arg
 Ala
 Arg
 Pro
 Thr
 Ile
 Lys
 Asn
 Lys
 His
 Gly
 Glu
 Ser

 Ser
 Leu
 Asp
 Ile
 Ala
 Arg
 Arg
 Leu
 Lys
 Phe
 Ser
 Gln
 Ile
 Glu
 Leu
 Met

Leu Arg Lys Ala Leu * 65 69

<210> 1151

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1151

<210> 1152

<211> 64

<212> PRT

<213> Homo sapiens

<400> 1152

<210> 1153

<211> 61

<212> PRT

<213> Homo sapiens

<400> 1153

<210> 1154 <211> 75

<212> PRT <213> Homo sapiens

<210> 1155 <211> 68 <212> PRT <213> Homo sapiens

| Met | Met | Ala | Lys | Ser | Val | Arg | Phe | Cys | Tyr | Val | Leu | Phe | Val | Glu | Glu | Glu | I | Ser | Ser | Val | Leu | Val | Glu | Glu | Glu | I | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Ser | Se

<210> 1156 <211> 60 <212> PRT <213> Homo sapiens

<210> 1157 <211> 776 <212> PRT

<213> Homo sapiens

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440
Ser Met Gln Leu Trp Gly Ser Thr Ser Asn Asp Gly Ala Phe Pro Ile
                   455
                                      460
Thr Asn Ile Ser Gly Thr Ser Met Gly Arg Tyr Ser Cys Cys Tyr His
                 470
                                   475
Pro Asp Trp Thr Ser Ser Ile Lys Ile Gln Pro Ser Asn Thr Leu Glu
           485
                               490
Leu Leu Val Thr Gly Leu Leu Pro Lys Pro Ser Leu Leu Ala Gln Pro
       500
                            505
Gly Pro Met Val Ala Pro Gly Glu Asn Met Thr Leu Gln Cys Gln Gly
                        520 525
Glu Leu Pro Asp Ser Thr Phe Val Leu Leu Lys Glu Gly Ala Gln Glu
                  535
Pro Leu Glu Gln Gln Arg Pro Ser Gly Tyr Arg Ala Asp Phe Trp Met
                 550
                                   555
Pro Ala Val Arg Gly Glu Asp Ser Gly Ile Tyr Ser Cys Val Tyr Tyr
              565
                               570
Leu Asp Ser Thr Pro Phe Ala Ala Ser Asn His Ser Asp Ser Leu Glu
                            585
Ile Trp Val Thr Asp Lys Pro Pro Lys Pro Ser Leu Ser Ala Trp Pro
                        600 605
Ser Thr Met Phe Lys Leu Gly Lys Asp Ile Thr Leu Gln Cys Arg Gly
                  615
                                      620
Pro Leu Pro Gly Val Glu Phe Val Leu Glu His Asp Gly Glu Glu Ala
                 630
                                   635
Pro Gln Gln Phe Ser Glu Asp Gly Asp Phe Val Ile Asn Asn Val Glu
             645
                               650 655
Gly Lys Gly Ile Gly Asn Tyr Ser Cys Ser Tyr Arg Leu Gln Ala Tyr
                         665
Pro Asp Ile Trp Ser Glu Pro Ser Asp Pro Leu Glu Leu Val Gly Ala
                        680
Ala Gly Pro Val Ala Gln Glu Cys Thr Val Gly Asn Ile Val Arg Ser
                  695
                                      700
Ser Leu Ile Val Val Val Val Ala Leu Gly Val Val Leu Ala Ile
     710
                                  715
Glu Trp Lys Lys Trp Pro Arg Leu Arg Thr Arg Gly Ser Glu Thr Asp
             725
                               730
Gly Arg Asp Gln Thr Ile Ala Leu Glu Glu Cys Asn Gln Glu Gly Glu
         740 745
Pro Gly Thr Pro Ala Asn Ser Pro Ser Ser Thr Ser Gln Arg Ile Ser
              760
Val Glu Leu Pro Val Pro Ile *
   770
                     775
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<210> 1158

<211> 80

<212> PRT

<213> Homo sapiens

<400> 1158

 Met
 Ile
 Gln
 Leu
 Phe
 Phe
 Val
 Leu
 Tyr
 Gly
 Ile
 Leu
 Ala
 Leu
 Ala
 Phe

 1
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 Leu
 Ser
 Gly
 Tyr
 Tyr
 Val
 Leu
 Ala
 Ala
 Gln
 Ile
 Leu
 Ala
 Val
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 Leu
 Ala
 Tyr
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 1
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 Tyr
 Trp
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Asn Thr Arg Arg Val Glu Phe Trp Asn Gln Met Lys Leu Leu Gly Glu
50 55 60

Ser Val Gly Ile Phe Gly Thr Ala Val Ile Leu Ala Thr Asp Gly *
65 70 75 79

<210> 1159 <211> 132 <212> PRT <213> Homo sapiens

<400> 1159 Met Ser Ser Gly Thr Glu Leu Leu Trp Pro Gly Ala Ala Leu Leu Val 10 Leu Leu Gly Val Ala Ala Ser Leu Cys Val Arg Cys Ser Arg Pro Gly 25 2.0 Ala Lys Arg Ser Glu Lys Ile Tyr Gln Gln Arg Ser Leu Arg Glu Asp 40 Gln Gln Ser Phe Thr Gly Ser Arg Thr Tyr Ser Leu Val Gly Gln Ala 55 Trp Pro Gly Pro Leu Ala Asp Met Ala Pro Thr Arg Lys Asp Lys Leu 75 70 Leu Gln Phe Tyr Pro Ser Leu Glu Asp Pro Ala Ser Ser Arg Tyr Gln 90 85 Asn Phe Ser Lys Gly Ser Arg His Gly Ser Glu Glu Ala Tyr Ile Asp 100 105 Pro Thr Ala Ile Lys Tyr Phe Leu Thr Gln Ala Thr Ala Ser Ile Ile 115 120 Leu Leu Ile Ala 130 132

<210> 1160 <211> 167 <212> PRT <213> Homo sapiens

<400> 1160 Met Val Gly Leu Gly Gly Met Ser Gln Leu Leu Leu Ala Ser Leu Leu Pro Pro Val Pro Gln Gly Ser Pro Thr Arg Arg Lys Leu Pro Ala Ser 25 Leu Leu Val Ser Thr Ala Leu Ile Ser Pro Val Cys Val Arg Gly Trp 40 Met Trp Gln Asn Leu Gln Asn Arg Ile His Gly Ser His Thr Ser Ala 55 Arg Arg Val Pro Ser Leu Pro Gly Ala Gly Gln Val Gly Val Arg Trp 70 75 Glu Ala Gly Pro Ala Cys Arg Thr Gln Pro Ser Pro Gln Asn Leu Ala 90 Pro Arg Pro His Pro Ser Ala Ala Gln Leu Ile Glu Asn Ala Ala Leu 105 Arg Ser Ala Met Ser Gly Glu Arg Leu Phe Pro Glu Gly Gln Glu His 120 Leu Gly Pro Leu Val Ala Pro Arg Val Pro Met Gly Gly Ala Leu Cys

<210> 1161 <211> 84 <212> PRT <213> Homo sapiens

<210> 1162 <211> 80 <212> PRT <213> Homo sapiens

<210> 1163 <211> 71 <212> PRT <213> Homo sapiens

<400> 1163
Met Tyr Gly Leu Lys Ile Leu Ser His Leu Trp Val Leu Leu Ile Leu
1 5 10 15

 Ser
 Leu
 Leu
 Phe
 Leu
 Arg
 Lys
 Ser
 Phe
 Lys
 Phe
 Tyr
 Ala
 Val
 Ser

 Phe
 Val
 Cys
 Phe
 Ala
 Phe
 Val
 Ala
 Phe
 Trp
 Asn
 Leu
 Gln
 Lys
 Ile

 Ile
 Ala
 Gln
 Ala
 Asn
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<210> 1164 <211> 56 <212> PRT <213> Homo sapiens

<210> 1165

<211> 97
<212> PRT
<213> Homo sapiens

<221> misc_feature
<222> (1)...(97)
<223> Xaa = any amino acid or nothing

<210> 1166 <211> 48

<212> PRT <213> Homo sapiens

<210> 1167 <211> 274 <212> PRT <213> Homo sapiens

<400> 1167 Met Glu Ala Pro Leu Ser His Leu Glu Ser Arg Tyr Leu Pro Ala His 1 5 Phe Ser Pro Leu Val Phe Phe Leu Leu Ser Ile Met Met Ala Cys 25 Cys Leu Val Ala Phe Phe Val Leu Gln Arg Gln Pro Arg Cys Trp Glu 40 Ala Ser Val Glu Asp Leu Leu Asn Asp Gln Val Thr Leu His Ser Ile Arg Pro Arg Glu Glu Asn Asp Leu Gly Pro Ala Gly Thr Val Asp Ser 75 Ser Gln Gly Gln Gly Tyr Leu Glu Glu Lys Ala Ala Pro Cys Cys Pro 90 Ala His Leu Ala Phe Ile Tyr Thr Leu Val Ala Phe Val Asn Ala Leu 105 Thr Asn Gly Met Leu Pro Ser Val Gln Thr Tyr Ser Cys Leu Ser Tyr 120 125 Gly Pro Val Ala Tyr His Leu Ala Ala Thr Leu Ser Ile Val Ala Asn 135 140 Pro Leu Ala Ser Leu Val Ser Met Phe Leu Pro Asn Arg Ser Leu Leu 150 155 Phe Leu Gly Val Leu Ser Val Leu Gly Thr Cys Phe Gly Gly Tyr Asn 165 170 Met Ala Met Ala Val Met Ser Pro Cys Pro Leu Leu Gln Gly His Trp 185 Gly Gly Glu Val Leu Ile Val Ser Ile Arg Pro Val Ala Ser Trp Val 200 Leu Phe Ser Gly Cys Leu Ser Tyr Val Lys Val Met Leu Gly Val Val 215 220 Leu Arg Asp Leu Ser Arg Ser Ala Leu Leu Trp Cys Gly Ala Ala Val 230 235 Gln Leu Gly Ser Leu Leu Gly Ala Leu Leu Met Phe Pro Leu Val Asn 250 Val Leu Arg Leu Phe Ser Ser Ala Asp Phe Cys Asn Leu His Cys Pro 265 Ala * 273

<210> 1168 <211> 230 <212> PRT <213> Homo sapiens

<400> 1168 Met Arg Ile Cys Asn Leu Ile Ser Met Met Leu Leu Cys His Trp 10 Asp Gly Cys Leu Gln Phe Leu Val Pro Met Leu Gln Asp Phe Pro Arg 20 25 Asn Cys Trp Val Ser Ile Asn Gly Met Val Asn His Ser Trp Ser Glu 40 Leu Tyr Ser Phe Ala Leu Phe Lys Ala Met Ser His Met Leu Cys Ile 55 Gly Tyr Gly Arg Gln Ala Pro Glu Ser Met Thr Asp Ile Trp Leu Thr Met Leu Ser Met Ile Val Gly Ala Thr Cys Tyr Ala Met Phe Ile Gly 90 His Ala Thr Ala Leu Ile Gln Ser Leu Asp Ser Ser Arg Arg Gln Tyr 105 Gln Glu Lys Tyr Lys Gln Val Glu Gln Tyr Met Ser Phe His Lys Leu 120 Pro Ala Asp Phe Arg Gln Lys Ile His Asp Tyr Tyr Glu His Arg Tyr 135 140 Gln Gly Lys Met Phe Asp Glu Asp Ser Ile Leu Gly Glu Leu Asn Gly 150 155 Pro Leu Arg Glu Glu Ile Val Asn Phe Asn Cys Arg Lys Leu Val Ala 170 165 Ser Met Pro Leu Phe Ala Asn Ala Asp Pro Asn Phe Val Thr Ala Met 185 180 Leu Thr Lys Leu Lys Phe Glu Val Phe Gln Pro Gly Asp Tyr Ile Ile 200 Pro Arg Arg His His Arg Glu Glu Asp Val Leu His Pro Ala Arg Arg 215 Gly Gln Arg Ala His * 229 225

<210> 1169 <211> 213 <212> PRT <213> Homo sapiens

85 90 Val Leu Met Ala Gly Ala Leu Ala Val Leu Ser Glu Gly Leu Gln Gly 105 Leu Asp Asp Glu Ala His Val Val Leu Ile Asp Val Glu Pro Gln Gln 120 125 Pro Gln Ala Ala Arg Gly Ala Ala Ala His Asp Val Gln Glu Leu Gln 135 140 Arg Leu Ala Tyr Gln Val Val Val Gly Phe Val Val Leu Thr Ala Gln 150 155 Glu Val Leu Gln Val Pro Val Val Val Leu Thr Gln Gln Leu Gln Lys 165 170 Ala Gln Asp Gly Leu His Asp Glu His Gly Cys Ala His Leu Thr Ala 180 Leu His Thr Phe Ala His Leu Val Pro Pro Ala Gln Ala Gly Ala Gln 195 200 Arg Val Ala Gly * 210 212

<210> 1170

<211> 51

<212> PRT

<213> Homo sapiens

<400> 1170

 Met
 Tyr
 Ser
 Leu
 Val
 Leu
 Thr
 Phe
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 Val
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 Phe
 Cys
 Ala
 Leu
 Ser

 Lys
 Thr
 Phe
 Leu
 Asp
 His
 Trp
 Phe
 Gln
 Met
 Phe
 Ile
 Tyr
 Tyr
 Ile
 Leu

 Phe
 Lys
 Asp
 Ser
 Glu
 Ile
 Gly
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 Pro
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 Tyr
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<210> 1171

<211> 157

<212> PRT

<213> Homo sapiens

<400> 1171

 Met
 Leu
 Val
 Pro
 Leu
 Asn
 Leu
 Cys
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 Ala
 Leu
 Val

 Ser
 Leu
 Pro
 Leu
 Pro
 Gly
 Ile
 Gly
 Arg
 Ala
 Phe
 Cys
 Glu
 Trp
 Leu
 Ser

 Gly
 Thr
 Phe
 Lys
 Ala
 Arg
 Arg
 Gln
 Gly
 Pro
 Lys
 Ala
 Lys
 Arg
 Glu
 Leu

 35
 Lys
 Ala
 Arg
 Arg
 Gly
 Pro
 Lys
 Ala
 Lys
 Arg
 Glu
 Leu

 Trp
 Asp
 Val
 Pro
 Val
 Arg
 Gly
 Trp
 Pro
 Trp
 Gly
 Phe
 Arg
 Leu

 Arg
 Gly
 Val
 Pro
 Val
 Ser
 Pro
 Ala
 Phe
 Gly
 Pro
 Phe
 Gly
 Ala

 Phe
 Gly
 <td

Arg Ser Ala Gly Arg Ser His Arg Gly Arg Arg Arg Arg Ala Ser Cys
115

Thr Ala Ala Pro Gly Gly Gly Val Thr Arg Arg Trp Lys Glu Tyr Cys
130

Thr Gln Arg Ile Asn Asn Leu Val Lys Pro Phe Ser *
145

<210> 1172 <211> 69 <212> PRT <213> Homo sapiens

<210> 1173 <211> 75 <212> PRT <213> Homo sapiens

<210> 1174 <211> 77 <212> PRT <213> Homo sapiens

<210> 1175 <211> 59 <212> PRT <213> Homo sapiens

<210> 1176 <211> 55 <212> PRT <213> Homo sapiens

<210> 1177 <211> 86 <212> PRT <213> Homo sapiens

 Ser
 Trp
 Val
 Arg
 Thr
 Ala
 Trp
 Met
 Leu
 Gly
 Ser
 Thr
 Ser
 Arg
 Thr
 Arg

 Gly
 Leu
 Ser
 Arg
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<210> 1178

<211> 189

<212> PRT

<213> Homo sapiens

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<210> 1179 <211> 55 <212> PRT <213> Homo sapiens

<210> 1180 <211> 81 <212> PRT <213> Homo sapiens

<210> 1181 <211> 69 <212> PRT <213> Homo sapiens

<210> 1182 <211> 430 <212> PRT <213> Homo sapiens

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Ile Arg Cys Leu Ala Gln Pro His Pro Gly Thr Gly Val Pro Arg Ala
Ala Ala Glu Leu Pro Leu Glu Ala Glu Lys Ile Lys Thr Gly Thr Gln
            100
                                105
                                                    110
Lys Gln Ala Lys Thr Asp Met Ala Phe Lys Thr Ser Val Ala Val Glu
                                                125
                            120
Met Ala Gly Ala Pro Ser Trp Thr Lys Val Ala Glu Glu Gly Asp Lys
                       135
                                            140
Pro Pro His Gly Pro Arg Cys Pro Asn His Ala Cys Gln Arg Leu Gly
                    150
                                        155
Gly Leu Ser Ala Pro Pro Trp Ala Lys Pro Glu Asp Arg Gln Thr Gln
                                    170
Pro Gln Pro His Gly His Val Pro Gly Lys Thr Thr Gln Gly Gly Pro
                                185
Cys Pro Ala Ala Cys Glu Val Gln Gly Met Leu Val Pro Pro Met Ala
                            200
Pro Thr Gly His Ser Thr Cys Asn Val Glu Ser Trp Gly Asp Asn Gly
                        215
                                            220
Ala Thr Arg Ala Gln Pro Ser Met Pro Gly Gln Ala Val Pro Cys Gln
                                        235
                    230
Glu Asp Thr Val Gly Ser Leu Leu Ala Ser Leu Cys Ala Glu Val Ala
                245
                                    250
Gly Val Leu Ala Ser Gln Glu Asp Leu Arg Thr Leu Leu Ala Lys Ala
                                265
Leu Ser Gln Gly Glu Val Trp Ala Ala Leu Asn Gln Ala Leu Ser Lys
                            280
                                                285
Glu Val Leu Gly Ala Thr Val Thr Lys Ala Leu Pro Gln Ser Met Leu
                        295
                                            300
Ser Met Ala Leu Val Lys Ala Leu Ser Trp Ser Glu Leu Arg Leu Thr
                    310
                                        315
Leu Ser Arg Ala Leu Ser Arg Gly Glu Leu Arg Ala Glu Leu Thr Lys
                                    330
Val Met Gln Gly Lys Leu Ala Glu Val Leu Ser Lys Ala Leu Thr Glu
            340
                                345
Glu Glu Trp Val Ala Leu Ser Gln Ala Leu Cys Gln Gly Glu Leu Gly
                            360
Ala Leu Leu Ser Gln Ser Trp Cys Arg Val Ala Leu Arg Thr Gly Thr
                        375
Ile Leu Pro Lys Ala Ala Ser Lys Ser Thr Gly Ser Gly Val Thr Lys
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                    390
Thr Pro Ala Leu Val Lys Val Ala Cys Arg Arg Ser Pro Ser Ala Ala
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Trp Gly Pro Ser Leu Gly Pro Val Arg Pro Gln Thr Ser Lys
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<210> 1183 <211> 53 <212> PRT

<213> Homo sapiens

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Ser Thr Ser Pro Pro Gly Ser Met Phe Phe Ser Ser Pro Pro Ser Arg
35 40 45

Gly Ile Pro Ala *
50 52

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<210> 1185 <211> 294 <212> PRT <213> Homo sapiens

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Gly Glu Leu Ala Gly Gln Glu Glu Glu Ala Leu Glu Gly Leu Glu 215 Val Met Asp Val Phe Leu Arg Phe Ser Gly Leu His Leu Phe Arg Ala 230 235 Val Glu Pro Gly Leu Val Gln Lys Phe Ser Leu Arg Asp Cys Ser Pro 250 245 Arg Leu Ser Glu Glu Leu Tyr His Arg Cys Arg Leu Ser Asn Leu Glu 260 265 Gly Leu Gly Gly Arg Ala Gln Leu Ala Met Ala Leu Phe Glu Gln Glu 275 280 285 Gln Ala Asn Ser Thr * 290 293

<210> 1186 <211> 57 <212> PRT <213> Homo sapiens

<210> 1187 <211> 191 <212> PRT <213> Homo sapiens

<400> 1187 Met Asp Leu Asp Asn Ala Lys Tyr Ser Leu Leu Gly Phe Ala Leu Phe 5 10 Trp Val Val Gly Phe Phe Val Cys Leu Phe Trp Phe Leu Val 20 25 Phe Leu Pro Trp Cys Lys Thr Val Glu Ser Cys Leu Phe Thr Gly Leu 40 Gly Ser Ile Glu Val Cys Val Ser Ser Val Arg Phe Leu Leu Arg Thr 55 60 Ile Cys Ile Phe Asn Asn Ser Thr Ser Ser Arg Pro Ser Arg Asn 70 75 Glu Arg Gly Leu Val Ser Ser Pro Glu Leu Ala Leu Glu Cys Val His 85 90 Leu Ala Ala His Gly Leu Val Ala Leu Arg Gly Leu Ile Gln Leu Pro 105 Leu Gln Leu Pro Ala Val Gly Val Asp Ala Leu Gly Leu Leu Cys 120 Leu Leu Gln Leu Pro Leu Glu Leu Leu Asp Pro Gly Ile Ala Phe Leu 135 Cys Leu Leu Val Leu Leu Gly His Leu Ala Leu Val Leu His Leu

<210> 1188 <211> 216 <212> PRT <213> Homo sapiens

<400> 1188 Met Ser Pro Pro Leu Leu Leu Pro Leu Leu Leu Leu Pro Leu 1 5 10 Leu Asn Val Glu Pro Ala Gly Ala Thr Leu Ile Arg Ile Pro Leu Arg 20 25 Gln Val His Pro Gly Arg Arg Thr Leu Asn Leu Leu Arg Gly Trp Gly 40 Lys Pro Ala Glu Leu Pro Lys Leu Gly Ala Pro Ser Pro Gly Asp Lys 55 Pro Ala Ser Val Pro Leu Ser Lys Phe Leu Asp Ala Gln Tyr Phe Gly 70 75 Glu Ile Gly Leu Gly Thr Pro Pro Gln Asn Phe Thr Val Ala Phe Asp 90 85 Thr Gly Ser Ser Asn Leu Trp Val Pro Ser Arg Arg Cys His Phe Phe 100 105 Ser Val Pro Cys Trp Phe His His Arg Phe Asn Pro Asn Ala Ser Ser 120 125 Ser Phe Lys Pro Ser Gly Thr Lys Phe Ala Ile Gln Tyr Gly Thr Gly 135 Arg Val Asp Gly Ile Leu Ser Glu Asp Lys Leu Thr Ile Gly Gly Ile 150 155 Lys Gly Ala Ser Val Ile Phe Gly Glu Ala Leu Trp Gly Ile Gln Pro 165 170 Gly Ser Ser Leu Phe Pro Ala Pro Met Gly Tyr Trp Gly Leu Gly Phe 180 185 Pro Ile Leu Val Leu Trp Glu Gly Ile Ser Ala Pro Ala Gly Cys Thr 195 · 200 Gly Gly Ala Gly Ala Ile Gly *

<210> 1189 <211> 176 <212> PRT <213> Homo sapiens

Ala Leu Ala Ala Ala Val Pro Ser Met Thr Gln Leu Leu Gly Asp Pro Gln Ala Gly Ile Arg Arg Asn Val Ala Ser Ala Leu Gly Asn Leu Gly 75 Rro Glu Gly Leu Gly Glu Glu Leu Leu Gln Cys Glu Val Pro Gln Arg 90 Leu Leu Glu Met Ala Cys Gly Asp Pro Gln Pro Asn Val Lys Glu Ala 105 Ala Leu Ile Ala Leu Arg Ser Leu Gln Gln Glu Pro Gly Ile His Gln 120 Val Leu Val Ser Leu Gly Ala Ser Glu Lys Leu Ser Leu Leu Ser Leu 135 140 Gly Asn Gln Ser Leu Pro His Ser Ser Pro Arg Pro Ala Ser Ala Lys 150 155 His Cys Arg Lys Leu Ile His Leu Leu Arg Pro Ala His Ser Met * 170 175

<210> 1190

<211> 58

<212> PRT

<213> Homo sapiens

<400> 1190

<210> 1191

<211> 88

<212> PRT

<213> Homo sapiens

<400> 1191

 Met
 Gly
 Ile
 Cys
 Leu
 Thr
 Trp
 Lys
 Pro
 Pro
 Thr
 Gly
 Val
 Ser
 Val
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 Leu
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 Met
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 Gly
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<210> 1192 <211> 136 <212> PRT <213> Homo sapiens

<400> 1192 Met Val Cys Leu Arg Leu Pro Gly Gly Ser Cys Met Ala Val Leu Thr 1 5 10 Val Thr Leu Met Val Leu Ser Ser Pro Leu Ala Leu Ala Gly Asp Thr 20 25 Arg Pro Arg Phe Leu Glu Tyr Ser Thr Ser Glu Cys His Phe Phe Asn 35 40 45 Gly Thr Glu Arg Val Arg Tyr Leu Asp Arg Tyr Phe His Asn Gln Glu 55 Glu Asn Val Arg Phe Asp Ser Asp Val Gly Glu Phe Arg Ala Val Thr 70 75 Glu Leu Gly Arg Pro Asp Ala Glu Tyr Trp Asn Ser Gln Lys Asp Leu 90 Leu Gly Thr Ala Arg Arg Thr Ser Trp Ser Arg Ser Gly Ala Gly Trp 105 Thr Thr Thr Ala Asp Thr Thr Thr Gly Leu Trp Arg Ala Ser Gln Cys 120 Ser Gly Glu Ser Ile Leu Arg *

<210> 1193 <211> 99 <212> PRT <213> Homo sapiens

<400> 1193 Met Leu Ala Ser Arg Gln Ala Cys Cys Pro Pro Val Ser Ser Leu Phe 10 Leu Pro Leu Ser Pro Thr Leu Ser Gly Phe Phe Thr Val Cys Ser Val 25 Ser His Leu His Val Pro Arg Gly Pro Ala Arg Leu Cys Pro Arg Met 40 Ser His Gly Ser Pro Ser Gly Leu Pro Ala Glu Pro Ser Glu His Gly 55 60 Cys Leu Leu Val Val Gly Leu Gln Gln Asn Cys Thr Arg Leu Thr Ser 70 75 Pro Ile Leu Ser Ser Arg Gly Leu Arg Val Gln Arg Arg Val Asn Leu 90 Ala Asp * 98

<210> 1194 <211> 50 <212> PRT <213> Homo sapiens

<400> 1194

 Met
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 Arg
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<210> 1195 <211> 58 <212> PRT <213> Homo sapiens

<210> 1196 <211> 132 <212> PRT <213> Homo sapiens

<400> 1196 Met Leu Pro Asn Ser Ser Ser Leu Trp Leu Val Met Arg Ile Leu Ile 10 Phe Cys Val Ile Pro Ala Gly Gly Val Leu Gly Ala Pro Thr Ala Ala 20 25 Gly Leu Arg Pro Thr Gly Asp Val Ala Leu Arg Arg Pro Ala Gly Ser Val Glu Pro Ser Gly Ser Arg Gly Leu Arg Ala Ser Val Cys Gln Arg 55 60 Leu Ser Met Phe Leu Ala His Phe Leu Arg Gly His Phe Leu Trp Trp 75 Ile Leu Asp Gly Gln Arg Leu Gly Phe Pro Leu Ser Leu Ala Thr Trp 90 Asn Arg Arg Lys Lys Ser Leu Gln His Leu Leu His Lys His Val Leu 105 Pro Val Arg Arg His Ala Gly Pro Cys Arg Gly Pro Gln Thr Thr Ala 115 120 Arg Gly Pro Arg 130 132

<210> 1197 <211> 64

<212> PRT <213> Homo sapiens

<210> 1198 <211> 53 <212> PRT <213> Homo sapiens

<210> 1199
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<212> PRT
<213> Homo sapiens

<221> misc_feature
<222> (1) ... (50)
<223> Xaa = any amino acid or nothing

<210> 1200 <211> 49 <212> PRT

<213> Homo sapiens

<210> 1201 <211> 46 <212> PRT <213> Homo sapiens

<210> 1202 <211> 332 <212> PRT <213> Homo sapiens

<400> 1202 Met Pro Leu Pro Trp Ser Leu Ala Leu Pro Leu Leu Ser Trp Val 10 Ala Gly Gly Phe Gly Asn Ala Ala Ser Ala Arg His His Gly Leu Leu 2.0 25 Ala Ser Ala Arg Gln Pro Gly Val Cys His Tyr Gly Thr Lys Leu Ala 40 Cys Cys Tyr Gly Trp Arg Arg Asn Ser Lys Gly Val Cys Glu Ala Thr 55 Cys Glu Pro Gly Cys Lys Phe Gly Glu Cys Val Gly Pro Asn Lys Cys 70 75 Arg Cys Phe Pro Gly Tyr Thr Gly Lys Thr Cys Ser Gln Asp Val Asn 90 Glu Cys Gly Met Lys Pro Arg Pro Cys Gln His Arg Cys Val Asn Thr 105 His Gly Ser Tyr Lys Cys Phe Cys Leu Ser Gly His Met Leu Met Pro 120 125 Asp Ala Thr Cys Val Asn Ser Arg Thr Cys Ala Met Ile Asn Cys Gln 135 140 Tyr Ser Cys Glu Asp Thr Glu Glu Gly Pro Gln Cys Leu Cys Pro Ser 150 155 Ser Gly Leu Arg Leu Ala Pro Asn Gly Arg Asp Cys Leu Asp Ile Asp

170 165 Glu Cys Ala Ser Gly Lys Val Ile Cys Pro Tyr Asn Arg Arg Cys Val 185 180 Asn Thr Phe Gly Ser Tyr Tyr Cys Lys Cys His Ile Gly Phe Glu Leu 200 205 Gln Tyr Ile Ser Gly Arg Tyr Asp Cys Ile Asp Ile Asn Glu Cys Thr 215 Met Asp Ser His Thr Cys Ser His His Ala Asn Cys Phe Asn Thr Gln 230 235 Gly Ser Phe Lys Cys Lys Cys Lys Gln Gly Tyr Lys Gly Asn Gly Leu 245 250 Arg Cys Ser Ala Ile Pro Glu Asn Ser Val Lys Glu Val Leu Arg Ala 265 Pro Gly Thr Ile Lys Asp Arg Ile Lys Lys Leu Leu Ala His Lys Asn 280 Ser Met Lys Lys Lys Ala Lys Ile Lys Asn Val Thr Pro Glu Pro Thr 300 295 Arg Thr Pro Thr Pro Lys Val Asn Leu Gln Pro Phe Asn Tyr Glu Glu 315 310 Ile Val Ser Arg Gly Gly Asn Ser His Gly Gly * 330 331

<210> 1203

<211> 825

<212> PRT

<213> Homo sapiens

<400> 1203

Met Ala Arg Leu Gly Asn Cys Ser Leu Thr Trp Ala Ala Leu Ile Ile 10 Leu Leu Pro Gly Ser Leu Glu Glu Cys Gly His Ile Ser Val Ser Ala Pro Ile Val His Leu Gly Asp Pro Ile Thr Ala Ser Cys Ile Ile Lys Gln Asn Cys Ser His Leu Asp Pro Glu Pro Gln Ile Leu Trp Arg Leu Gly Ala Glu Leu Gln Pro Gly Gly Arg Gln Gln Arg Leu Ser Asp Gly Thr Gln Glu Ser Ile Ile Thr Leu Pro His Leu Asn His Thr Gln Ala Phe Leu Ser Cys Cys Leu Asn Trp Gly Asn Ser Leu Gln Ile Leu 105 Asp Gln Val Glu Leu Arg Ala Gly Tyr Pro Pro Ala Ile Pro His Asn 125 120 Leu Ser Cys Leu Met Asn Leu Thr Thr Ser Ser Leu Ile Cys Gln Trp 135 140 Glu Pro Gly Pro Glu Thr His Leu Pro Thr Ser Phe Thr Leu Lys Ser 155 150 Phe Lys Ser Arg Gly Asn Cys Gln Thr Gln Gly Asp Ser Ile Leu Asp 165 170 Cys Val Pro Lys Asp Gly Gln Ser His Cys Cys Ile Pro Arg Lys His 185 Leu Leu Leu Tyr Gln Asn Met Gly Ile Trp Val Gln Ala Glu Asn Ala 200 Leu Gly Thr Ser Met Ser Pro Gln Leu Cys Leu Asp Pro Met Asp Val 215

Val 225	Lys	Leu	Glu	Pro	Pro 230	Met	Leu	Arg	Thr	Met 235	Asp	Pro	Ser	Pro	Glu 240
Ala	Ala	Pro	Pro	Gln 245	Ala	Gly	Cys	Leu	Gln 250	Leu	Cys	Trp	Glu	Pro 255	Trp
Gln	Pro	Gly	Leu 260	His	Ile	Asn	Gln	Lys 265	Cys	Glu	Leu	Arg	His 270	Lys	Pro
Gln	Arg	Gly 275	Glu	Ala	Ser	Trp	Ala 280	_	Val	Gly	Pro	Leu 285		Leu	Glu
Ala	Leu 290	_	Tyr	Glu	Leu	Cys 295		Leu	Leu	Pro	Ala 300		Ala	Tyr	Thr
Leu 305	Gln	Ile	Arg	Cys	Ile 310		Trp	Pro	Leu	Pro 315		His	Trp	Ser	Asp
Trp	Ser	Pro	Ser	Leu 325	Glu	Leu	Arg	Thr	Thr 330		Arg	Ala	Pro	Thr	
Arg	Leu	Asp	Thr 340	Trp	Trp	Arg	Gln	Arg 345		Leu	Asp	Pro	Arg 350	Thr	Val
Gln	Leu	Phe 355	Trp	Lys	Pro	Val	Pro 360	Leu	Glu	Glu	Asp	Ser 365	Gly	Arg	Ile
Gln	Gly 370	Tyr	Val	Val	Ser	Trp 375	Arg	Pro	Ser	Gly	Gln 380	Ala	Gly	Ala	Ile
Leu 385	Pro	Leu	Cys	Asn	Thr 390	Thr	Glu	Leu	Ser	Cys 395	Thr	Phe	His	Leu	Pro 400
Ser	Glu	Ala	Gln	Glu 405	Val	Ala	Leu	Val	Ala 410	Tyr	Asn	Ser	Ala	Gly 415	Thr
Ser	Arg	Pro	Thr 420	Pro	Val	Val	Phe	Ser 425	Glu	Ser	Arg	Gly	Pro 430	Ala	Leu
Thr	Arg	Leu 435	His	Ala	Met	Ala	Arg 440	Asp	Pro	His	Ser	Leu 445	Trp	Val	Gly
	Glu 450					455					460			_	_
Leu 465	Gly	Pro	Pro	Ser	Ala 470	Ser	Asn	Ser	Asn	Lys 475	Thr	Trp	Arg	Met	Glu 480
Gln	Asn	Gly	Arg	Ala 485	Thr	Gly	Phe	Leu	Leu 490	ГЛЗ	Glu	Asn	Ile	Arg 495	Pro
	Gln		500					505			_		510		
	Pro	515					520					525			
	Ala 530					535					540				
545					550					555					560
	Tyr			565					570					575	
	Leu		580					585					590		
	Ser	595					600					605		_	
	Asn 610					615			•		620				
Gly 625	Arg	Ile	Pro	Ser	Gly 630	Gln	Val	Ser	Gln	Thr 635	Gln	Leu	Thr	Ala	Ala 640
	Ala	Pro	Gly	Cys 645		Gln	Ser	Trp	Arg 650		Met	Pro	Ser	Ser 655	
Pro	Ala	Leu	Ala 660		His	Pro	Ser	Pro 665		Ser	Gln	Cys	Trp 670		Arg
Met	Lys	Arg 675		Arg	Cys	Pro	Gly 680		Pro	Ile	Thr	Ala 685		Arg	Pro
Val	Ala	Ser	Pro	Leu	Trp	Ser	Arg	Pro	Met	Cys	Ser	Arg	Gly	Thr	Gln

695 Glu Gln Phe Pro Pro Ser Pro Asn Pro Ser Leu Ala Pro Ala Ile Arg 710 715 720 Ser Phe Met Gly Ser Cys Trp Ala Ala Pro Gln Ala Gln Gly Gln Gly 725 730 Thr Ile Ser Ala Val Thr Pro Leu Ser Pro Ser Trp Arg Ala Ser Pro 740 745 Pro Ala Pro Ser Pro Met Arg Thr Ser Gly Ser Arg Pro Ala Pro Trp 760 Gly Pro Leu Val Thr Pro Ser Pro Lys Ser Gln Glu Asp Asp Cys Val 775 Phe Gly Pro Leu Leu Asn Phe Pro Pro Ser Cys Arg Gly Ser Gly Ser 790 795 Met Gly Trp Arg Arg Trp Gly Ala Ser Arg Ala Ser Leu Gly Phe Pro 805 810 Ser Trp Ala Cys Leu Leu Lys Ala * 820

<210> 1204

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1204

<210> 1205

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1205

<210> 1206

<211> 88

<212> PRT

<213> Homo sapiens

<400> 1206

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 Glu
 Ala

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 Gly
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<210> 1207 <211> 186 <212> PRT <213> Homo sapiens

<400> 1207 Met Ile Leu Asn Lys Ala Leu Met Leu Gly Ala Leu Ala Leu Thr Thr Val Met Ser Pro Cys Gly Gly Glu Asp Ile Val Ala Asp His Val Ala 2.0 25 Ser Tyr Gly Val Asn Leu Tyr Gln Ser Tyr Gly Pro Ser Gly Gln Tyr 40 Ser His Glu Phe Asp Gly Asp Glu Glu Phe Týr Val Asp Leu Glu Arg 55 Lys Glu Thr Val Trp Gln Leu Pro Leu Phe Arg Arg Phe Arg Arg Phe 70 75 Asp Pro Gln Phe Ala Leu Thr Asn Ile Ala Val Leu Lys His Asn Leu 90 85 Asn Ile Val Ile Lys Arg Ser Asn Ser Thr Ala Ala Thr Asn Glu Val 105 Pro Glu Val Thr Val Phe Ser Lys Ser Pro Val Thr Leu Gly Gln Pro 120 Asn Thr Leu Ile Cys Leu Val Asp Asn Ile Phe Pro Pro Val Val Asn 140 135 Ile Thr Trp Leu Ser Asn Gly His Ser Val Thr Glu Gly Val Ser Glu 155 150 Thr Arg Pro Ser Ser Pro Lys Ser Asp His Phe Leu Leu Gln Asp Gln 165 170 175 Val Thr Ser Pro Ser Phe Pro Phe Glu * 180 185

<210> 1208 <211> 46 <212> PRT <213> Homo sapiens

20 25 30
Pro Ser Ser Arg Met Trp Lys Ser Ile Ile Phe Phe Leu *
35 40 45

<210> 1209 <211> 199 <212> PRT <213> Homo sapiens

<400> 1209 Met Ala Leu Leu Val Pro Leu Ala Leu Leu Val Ile Gln Ala His Leu 1 5 10 Val Leu Ser Val Gln Leu Glu Arg Val Val Thr Glu Glu Lys Val Ala 25 Leu Leu Ala Leu Leu Val Leu Pro Val Leu Leu Val Pro Glu Val Leu 40 Leu Val Leu Lys Ala His Val Val Thr Lys Val Lys Gln Val Asn Val 55 Glu Leu Leu Ala Ser Lys Asp Ile Glu Asp Ser Leu Val Ile Gln Val 70 75 Pro Gln Val Leu Gln Ala Leu Leu Val Ser Arg Val Gln Ser Ala Val 90 85 Gln Asp Leu Gln Ala Pro Glu Asp Leu Leu Asp Pro Val Asp Leu Leu 105 100 Ala Lys Met Glu Pro Val Asp Ile Gln Val Pro Leu Asp His Gln Gly 120 125 Leu Glu Val Thr Glu Val Lys Glu Asp Leu Arg Ala Pro Gln Ala Thr 135 140 Gln Gly Asn Gln Ala Leu Leu Asp Leu Leu Val Pro Leu Val Leu Ala 150 155 Val Val Leu Glu Pro Leu Pro Leu Gly Leu Glu Val Lys Lys 165 170 Leu Ala Val Leu Pro Arg Ile Met Glu Met Asn Gln Trp Ile Ser Lys 180 185 Ser Thr Pro Met Arg Leu * 195 198

<210> 1210 <211> 59 <212> PRT <213> Homo sapiens

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<210> 1211
<211> 227
<212> PRT
<213> Homo sapiens

<221> misc_feature
<222> (1)...(227)
<223> Xaa = any amino acid or nothing

<400> 1211
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Met Ala Ser Ile Cys Ser Trp Arg Val Met Leu Ala Trp Ala Ala Cys Trp Val Arg Ala His Ala Ala Leu Ser Gly His Pro Arg Ser Thr Phe 25 Ser Leu Trp Leu Ser Gly Ile Ser Leu Pro Xaa Pro Ile Phe Leu Pro 40 4.5 Met Ala Val Ser Leu Leu Thr Pro Lys Asp Val Lys Tyr Ala Arg Ser 55 Pro Asn Cys Phe Lys Ala Ala Leu Asn Ile Pro Asp Pro Gly Ala Val 70 75 His Leu Ile Ile Ala Leu Leu Thr Asp Gly Ala Ile Pro Leu Leu 90 85 Gln Pro Ala Arg Val Lys Lys Ser Asn Ala His Val Phe Leu His Phe 105 Ala Gly Gly Asp Leu Leu Pro Ser Asn Gly Gly His Lys Ile Leu Ile 120 125 Trp Ser Arg Gly Trp Arg Gln Gly Leu Gly Gly Phe Gly Ile Ile Ile 135 Leu Ala Asp Asn Asp Leu Val Trp Ser Trp Gly Gln Ser Trp Arg His 150 155 Gly Cys Leu Leu Gly Val Gly Ala Leu Ser Ala Leu Leu Leu His His 165 170 Leu Asn Pro His Pro Tyr Leu Val Leu Gly Cys Pro Gly Pro Ala Gly 180 185 Lys Glu Ala Pro Pro Pro Ser Pro Val Cys His Pro Pro His Gln Thr 200 Arg Pro Pro Ser Gln Leu Pro His Ser Pro Gln Thr Phe His Ser Ala 215 Pro Glu 225 226

<210> 1212 <211> 62 <212> PRT <213> Homo sapiens

50 55 60 61

<210> 1213 <211> 55 <212> PRT

<213> Homo sapiens

<210> 1214 <211> 642 <212> PRT <213> Homo sapiens

<400> 1214 Met Thr Met Tyr Leu Trp Leu Lys Leu Leu Ala Phe Gly Phe Ala Phe Leu Asp Thr Glu Val Phe Val Thr Gly Gln Ser Pro Thr Pro Ser Pro 25 Thr Asp Ala Tyr Leu Asn Ala Ser Glu Thr Thr Leu Ser Pro Ser 40 Gly Ser Ala Val Ile Ser Thr Thr Thr Ile Ala Thr Thr Pro Ser Lys 55 Pro Thr Cys Asp Glu Lys Tyr Ala Asn Ile Thr Val Asp Tyr Leu Tyr 70 75 Asn Lys Glu Thr Lys Leu Phe Thr Ala Lys Leu Asn Val Asn Glu Asn 90 Val Glu Cys Gly Asn Asn Thr Cys Thr Asn Asn Glu Val His Asn Leu 105 Thr Glu Cys Lys Asn Ala Ser Val Ser Ile Ser His Asn Ser Cys Thr 120 115 Ala Pro Asp Lys Thr Leu Ile Leu Asp Val Pro Pro Gly Val Glu Lys 135 140 Phe Gln Leu His Asp Cys Thr Gln Val Glu Lys Ala Asp Thr Thr Ile 150 155 Cys Leu Lys Trp Lys Asn Ile Glu Thr Phe Thr Cys Asp Thr Gln Asn 170 Ile Thr Tyr Arg Phe Gln Cys Gly Asn Met Ile Phe Asp Asn Lys Glu 180 185 Ile Lys Leu Glu Asn Leu Glu Pro Glu His Glu Tyr Lys Cys Asp Ser 200 205 Glu Ile Leu Tyr Asn Asn His Lys Phe Thr Asn Ala Ser Lys Ile Ile 215 220 Lys Thr Asp Phe Gly Ser Pro Gly Glu Pro Gln Ile Ile Phe Cys Arg 230 235

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Ser Glu Ala Ala His Gln Gly Val Ile Thr Trp Asn Pro Pro Gln Arg
               245
                                   250
Ser Phe His Asn Phe Thr Leu Cys Tyr Ile Lys Glu Thr Glu Lys Asp
           260
                               265
Cys Leu Asn Leu Asp Lys Asn Leu Ile Lys Tyr Asp Leu Gln Asn Leu
      275
                           280
Lys Pro Tyr Thr Lys Tyr Val Leu Ser Leu His Ala Tyr Ile Ile Ala
                       295
                                           300
Lys Val Gln Arg Asn Gly Ser Ala Ala Met Cys His Phe Thr Thr Lys
                   310
                                       315
Ser Ala Pro Pro Ser Gln Val Trp Asn Met Thr Val Ser Met Thr Ser
                                        335
                                   330
Asp Asn Ser Met His Val Lys Cys Arg Pro Pro Arg Asp Arg Asn Gly
                               345
Pro His Glu Arg Tyr His Leu Glu Val Glu Ala Gly Asn Thr Leu Val
                           360
Arg Asn Glu Ser His Lys Asn Cys Asp Phe Arg Val Lys Asp Leu Gln
                       375
Tyr Ser Thr Asp Tyr Thr Phe Lys Ala Tyr Phe His Asn Gly Asp Tyr
                                       395
                   390
Pro Gly Glu Pro Phe Ile Leu His His Ser Thr Ser Tyr Asn Ser Lys
                405
                                   410
Ala Leu Ile Ala Phe Leu Ala Phe Leu Ile Ile Val Thr Ser Ile Ala
                                425
            420
Leu Leu Val Val Leu Tyr Lys Ile Tyr Asp Leu His Lys Lys Arg Ser
                           440
Cys Asn Leu Asp Glu Gln Glu Leu Val Glu Arg Asp Asp Glu Lys
                       455
                                           460
Gln Leu Met Asn Val Glu Pro Ile His Ala Asp Ile Leu Leu Glu Thr
                                       475
                    470
Tyr Lys Arg Lys Ile Ala Asp Glu Gly Arg Leu Phe Leu Ala Glu Phe
                485
                                   490
Gln Ser Ile Pro Arg Val Phe Ser Lys Phe Pro Ile Lys Glu Ala Arg
                               505
            500
Lys Pro Phe Asn Gln Asn Lys Asn Arg Tyr Val Asp Ile Leu Pro Tyr
                           520
Asp Tyr Asn Arq Val Glu Leu Ser Glu Ile Asn Gly Asp Ala Gly Ser
                        535
                                           540
Asn Tyr Ile Asn Ala Ser Tyr Ile Asp Gly Phe Lys Glu Pro Arg Lys
                                       555
Tyr Ile Ala Ala Gln Gly Pro Arg Asp Glu Thr Val Asp Asp Phe Trp
                                    570
Arg Met Ile Trp Glu Gln Lys Ala Thr Val Ile Val Met Val Thr Arg
            580
                                585
                                                   590
Cys Glu Glu Gly Asn Arg Asn Lys Cys Ala Glu Tyr Trp Pro Ser Met
                           600
Glu Glu Gly Thr Arg Ala Phe Gly Glu Cys Cys Cys Lys Asp Leu Thr
                       615
                                           620
Lys His Lys Arg Cys Pro Arg Leu His His Ser Glu Ile Glu His Cys
625
                   630
                                       635
Lys *
641
```

<210> 1215

<211> 85

<212> PRT

<213> Homo sapiens

<210> 1216 <211> 403 <212> PRT <213> Homo sapiens

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Pro Gln Pro Val Met Val Thr Trp Val Arg Val Asp Asp Glu Met Pro 280 275 Gln His Ala Val Leu Ser Gly Pro Asn Leu Phe Ile Asn Asn Leu Asn 295 300 Lys Thr Asp Asn Gly Thr Tyr Arg Cys Glu Ala Ser Asn Ile Val Gly 310 315 Lys Ala His Ser Asp Tyr Met Leu Tyr Val Tyr Asp Pro Pro Thr Thr 325 330 345 Thr Ile Leu Thr Ile Ile Thr Asp Ser Arg Ala Gly Glu Gly Ser 360 365 Ile Arg Ala Val Asp His Ala Val Ile Gly Gly Val Val Ala Val Val 375 380 Val Phe Ala Met Leu Cys Leu Leu Ile Ile Leu Gly Arg Tyr Phe Ala 390 395 Gln Thr * 402

<210> 1217 <211> 49 <212> PRT

<213> Homo sapiens

<210> 1218 <211> 304 <212> PRT <213> Homo sapiens

<400> 1218 Met Ala Arg Arg Ser Arg His Arg Leu Leu Leu Leu Leu Arg Tyr 10 Leu Val Val Ala Leu Gly Tyr His Lys Ala Tyr Gly Phe Ser Ala Pro Lys Asp Gln Gln Val Val Thr Ala Val Glu Tyr Gln Glu Ala Ile Leu 40 Ala Cys Lys Thr Pro Lys Lys Thr Val Ser Ser Arg Leu Glu Trp Lys 55 60 Lys Leu Gly Arg Ser Val Ser Phe Val Tyr Tyr Gln Gln Thr Leu Gln 70 75 Gly Asp Phe Lys Asn Arg Ala Glu Met Ile Asp Phe Asn Ile Arg Ile 85 90 Lys Asn Val Thr Arg Ser Asp Ala Gly Lys Tyr Arg Cys Glu Val Ser

			100					105					110		
Ala	Pro	Ser	Glu	Gln	Gly	Gln	Asn 120	Leu	Glu	Glu	Asp	Thr 125	Val	Thr	Leu
Glu	Val 130	Leu	Gly	Asp	Val	His 135	Val	Leu	Ala	Pro	Ala 140	Val	Pro	Ser	Cys
Glu 145	Val	Pro	Ser	Ser	Ala 150	Leu	Ser	Gly	Thr	Val 155	Val	Glu	Leu	Arg	Cys 160
Gln	Asp	Lys	Glu	Gly 165	Asn	Pro	Ala	Pro	Glu 170	Tyr	Thr	Trp	Phe	Lys 175	Asp
Gly	Ile	Arg	Leu 180	Leu	Glu	Asn	Pro	Arg 185		Gly	Ser	Gln		Thr	Asn
Ser	Ser	Tyr 195	Thr	Met	Asn	Thr	Lys 200	Thr	Gly	Thr	Leu	Gln 205	Phe	Asn	Thr
Val	Ser 210	Lys	Leu	Asp	Thr		Glu		Ser	Cys	Glu 220	Ala	Arg	Asn	Ser
Val 225	Gly	Tyr	Arg	Arg	Cys 230	Pro	Gly	Lys	Arg	Met 235	Gln	Val	Asp	Asp	Leu 240
Asn	Ile	Ser	Gly	Ile 245	Ile	Ala	Ala	Val	Val 250	Val	Val	Ala	Leu	Val 255	Ile
Ser	Val	Cys	Gly 260	Leu	Gly	Val	Cys	Tyr 265	Ala	Gln	Arg	Lys	Gly 270	Tyr	Phe
Ser	Lys	Glu 275	Thr	Ser	Phe	Gln	Lys 280	Ser	Asn	Ser	Ser	Ser 285	Lys	Ala	Thr
Thr	Met 290	Ser	Glu	Asn	Asp	Phe 295	Lys	His	Thr	Lys		Phe		Ile 303	*

<210> 1219

<211> 1126

<212> PRT

<213> Homo sapiens

<400> 1219

Met Trp Phe Leu Phe Leu Cys Pro Asn Leu Trp Ala Met Pro Val Gln 1 5 10 Ile Ile Met Gly Val Ile Leu Leu Tyr Asn Leu Leu Gly Ser Ser Ala 20 25 Leu Val Gly Ala Ala Val Ile Val Leu Leu Ala Pro Ile Gln Tyr Phe 40 Ile Ala Thr Lys Leu Ala Glu Ala Gln Lys Ser Thr Leu Asp Tyr Ser 55 Thr Glu Arg Leu Lys Lys Thr Asn Glu Ile Leu Lys Gly Ile Lys Leu 70 Leu Lys Leu Tyr Ala Trp Glu His Ile Phe Cys Lys Ser Val Glu Glu 90 Thr Arg Met Lys Glu Leu Ser Ser Leu Lys Thr Phe Ala Leu Tyr Thr 105 Ser Leu Ser Ile Phe Met Asn Ala Ala Ile Pro Ile Ala Ala Val Leu 120 Ala Thr Phe Val Thr His Ala Tyr Ala Ser Gly Asn Asn Leu Lys Pro 135 140 Ala Glu Ala Phe Ala Ser Leu Ser Leu Phe His Ile Leu Val Thr Pro 150 155 Leu Phe Leu Leu Ser Thr Val Val Arg Phe Ala Val Lys Ala Ile Ile 165 170 Ser Val Gln Lys Leu Asn Glu Phe Leu Leu Ser Asp Glu Ile Gly Asp 180 185

Asp	Ser	Trp 195	Arg	Thr	Gly	Glu	Ser 200	Ser	Leu	Pro	Phe	Glu 205	Ser	Cys	Lys
Lys	His 210	Thr	Gly	Val	Gln	Pro 215	Lys	Thr	Ile	Asn	Arg 220	Lys	Gln	Pro	Gly
Ara	Tyr	His	Ten	Asp	Ser		Glu	Gln	Ser	Thr		Ara	Leu	Ara	Pro
225	- 7 -	1110	LCu	прр	230	- y -	024	0.111	501	235	9	1119			240
	C7.11	Tha	<i>α</i> 1	7 ~~	_	777	T10	Tara	7707		7.00	Clar	T'3 22	Dho	
Ата	Glu	TIIL	Gru		116	нта	116	цуѕ		TIIT	ASII	GIY	TYL		Ser
_	~3	_	~-7	245	~ 7 .		-	_	250			-1-		255	
Trp	Gly	ser	_	Leu	Ата	Thr	ьeu		Asn	TTE	Asp	TTE		тте	Pro
_	_	_	260	_		_	_	265	_	_	_		270		
Thr	Gly	Gln	Leu	Thr	Met	Ile		Gly	Gln	Val	Gly	_	Gly	Lys	Ser
		275					280					285			
Ser	Leu	Leu	Leu	Ala	Ile	Leu	Gly	Glu	Met	Gln	Thr	Leu	Glu	Gly	Lys
	290					295					300				
Val	His	Trp	Ser	Asn	Val	Asn	Glu	Ser	Glu	Pro	Ser	Phe	Glu	Ala	Thr
305					310					315					320
Arg	Ser	Arg	Asn	Arg	Tyr	Ser	Val	Ala	Tyr	Ala	Ala	Gln	Lys	Pro	\mathtt{Trp}
				325					330					335	
Leu	Leu	Asn	Ala	Thr	Val	Glu	Glu	Asn	Ile	Thr	Phe	Gly	Ser	Pro	Phe
			340					345					350		
Asn	Lys	Gln	Arg	Tyr	Lys	Ala	Val	Thr	Asp	Ala	Cys	Ser	Leu	Gln	Pro
		355					360					365			
Asp	Ile	Asp	Leu	Leu	Pro	Phe	Gly	Asp	Gln	Thr	Glu	Ile	Gly	Glu	Arg
_	370	_				375	_	-			380		_		_
Gly	Ile	Asn	Leu	Ser	Gly	Gly	Gln	Arq	Gln	Arg	Ile	Cys	Val	Ala	Arq
385					390	-		_		395		-			400
Ala	Leu	Tyr	Gln	Asn	Thr	Asn	Ile	Val	Phe	Leu	Asp	Asp	Pro	Phe	Ser
		- 1 -		405					410		1	1		415	
Ala	Leu	Asp	Ile		Leu	Ser	Asp	His		Met	Gln	Glu	Glv		Leu
			420					425					430		
Lvs	Phe	Len		Asp	Asp	Lvs	Ara		Leu	Val	Len	Val		His	Tivs
1	- 110	435	0111	TLOP	Tabp	ביים	440		Dea	van		445		1110	2,2
T.e11	Gln		Len	Thr	His	Δla		Tro	Tle	Tle	Δla		Lvs	Asn	Glv
Lou	450	- 1 -			113.0	455	p	115			460		-7-	1101	017
Ser	Val	Tien	Ara	Glu	Glv		Ten	Lvs	Asp	Tle		Thr	Tvs	Asp	Val
465	var	cu	****9	<u></u>	470		Lea	цуз	TIDE	475	0111	1111	حا برحد	Tup	480
	Leu	ጥኒፖ	Glu	ніс		Taze	Thr	T.611	Met		Ara	Gln	Δen	Gln	
014	Dea	- y -	01.4	485	110	цуз	1111	пси	490	71011	Arg	3111	rup	495	GIG
T.011	Glu	Tare	7 cn		G111	λla	Λen	Gln		Thr	T.011	Glu	Δνα		Thr
пеп	Giu	цуѕ	500	MEC	GIU	Ата	Asp	505	1111	1111	пец	GIU	510	пув	1111
T. 011	Arg	71 200		Mot	Ma rae	C.~	7.~~		777	Taro	777-	~1 _~		C1	7 an
пеа	ALG	515		Mec	тут		520		Ата	цуъ	Ала	525	1.16.0	GIU	Asp
~7.,,	Asp			C1	<i>α</i> 3				7 00	C7.11	7.00		7 cn	Mot	Cox
GIU	530	GIU	Giu	Giu	GIU	535	GIU	Gru	Asp	Giu	540	ASP	ASII	Mer	per
The		Ma+	7\ ~~~	Τ	7 ~~~		T	740+	Dwo	Tren		mb~	Cara	There	71
	Val	Mec	Arg	ьеu		TILE	ьуѕ	Mer	PIO		ьуѕ	1111	Cys	тр	
545	T	m1	0	07	550	701	ml	T	T	555	Ŧ	34-4	~ 7~	Di	560
Tyr	Leu	Thr	ser	_	GTY	Pne	Pne	ьeu		тте	ьeu	мес	тте		ser
-	-		T	565		7		7	570	-1 -	_		m	575	27.
гув	Leu	ьeu		Hls	ser	vaı	тте		Ara	тте	Asp	Tyr		ьeu	Ата
	_		580			_		585	_			_	590	_	
Thr	\mathtt{Trp}		Ser	GLu	Tyr	Ser		Asn	Asn	Thr	GΤΛ	_	Ala	Asp	Gln
		595					600		_	_		605			
Thr	Tyr	Tyr	Val	Ala	Gly	Phe	Ser	Ile	Leu	Cys	Gly	Ala	Gly	Ile	Phe
	610					615					620				
	Cys	Leu	Val	Thr	Ser	Leu	Thr	Val	Glu	\mathtt{Trp}	Met	Gly	Leu	Thr	Ala
625					630					635					640
Ala	Lys	Asn	Leu	His	His	Asn	Leu	Leu		Lys	Ile	Ile	Leu	Gly	Pro
				645					650					655	
Ile	Arg	Phe	Phe	Asp	Thr	Thr	Pro	Leu	Gly	Leu	Ile	Leu	Asn	Arg	Phe

			660					665					670		
Ser	Ala	Asp 675	Thr	Asn	Ile	Ile	Asp 680	Gln	His	Ile	Pro	Pro 685		Leu	Glu
Ser	Leu 690	Thr	Arg	Ser	Thr	Leu 695	Leu	Cys	Leu	Ser	Ala 700	Ile	Gly	Met	Ile
Ser 705		Ala	Thr	Pro	Val 710	Phe	Leu	Val	Ala	Leu 715	Leu	Pro	Leu	Gly	Val 720
Ala	Phe	Tyr	Phe	Ile 725	Gln	Lys	Tyr	Phe	Arg 730	Val	Ala	Ser	Lys	Asp 735	Leu
Gln	Glu	Leu	Asp 740	Asp	Ser	Thr	Gln	Leu 745	Pro	Leu	Leu	Cys	His 750	Phe	Ser
Glu	Thr	Ala 755	Glu	Gly	Leu	Thr	Thr 760	Ile	Arg	Ala	Phe	Arg 765	His	Glu	Thr
Arg	Phe 770	Lys	Gln	Arg	Met	Leu 775	Glu	Leu	Thr	Asp	Thr 780	Asn	Asn	Ile	Ala
785					790				Trp	795					800
				805					Ala 810					815	
			820					825	Leu				830		
		835					840		Val			845		_	
	850					855			Val		860				
865					870				Asp	875					880
				885					Ile 890				_	895	_
			900					905	Lys			_	910		
		915					920		Gly			925		-	-
	930					935			Met		940			_	=
945					950				Ser	955					960
				965					Gln 970					975	
			980					985	Glu	_		_	990	_	_
		995				1	1000		Gln		1	.005			_
]	L010				1	.015			Val	1	.020	_	-		
1025	ser	val	GIY		Arg .030	GIn	Leu	Pne	Cys	ьеи 1035	Ala	Arg	Ala		Val .040
Arg	Lys	Ser		Ile .045	Leu	Ile	Met		Glu L050	Ala	Thr	Ala		Ile .055	Asp
		3	.060				ב	.065	Val			1	.070		
Asp		Thr 1075	Val	Val	Thr		Ala .080	His	Arg	Val		Ser .085	Ile	Met	Asp
]	1090				1	.095			Gly	1	100			_	-
Thr 1105	Val	Pro	Asn		Phe 110	Ala	His	Lys	Asn 1	Gly 115	Pro	Phe	Ser		Leu .120
Val	Met	Thr		Lys .125	*										

<210> 1220 <211> 46 <212> PRT <213> Homo sapiens

<210> 1221 <211> 56 <212> PRT <213> Homo sapiens

<210> 1222 <211> 253 <212> PRT <213> Homo sapiens

 <400> 1222

 Met Gly Cys Ala Ile Ile Ala Gly Phe Leu His Tyr Leu Phe Leu Ala 1

 5
 10
 15

 Cys Phe Phe Phe Trp Met Leu Val Glu Ala Val Ile Leu Phe Leu Met Val 20

 25
 30

 Arg Asn Leu Lys Val Val Asn Tyr Phe Ser Ser Arg Asn Ile Lys Met 35

 40
 45

 Leu His Ile Cys Ala Phe Gly Tyr Gly Leu Pro Met Leu Val Val Val 50

 55
 60

 Ile Ser Ala Ser Val Gln Pro Gln Gly Tyr Gly Met His Asn Arg Cys 65

 75
 80

 Trp Leu Asn Thr Glu Thr Gly Phe Ile Trp Ser Phe Leu Gly Pro Val 85

 90
 95

 Cys Thr Val Ile Val Ile Asn Ser Leu Leu Leu Thr Trp Thr Leu Trp 100

 105
 110

 Ile Leu Arg Gln Arg Leu Ser Ser Val Asn Ala Glu Val Ser Thr Leu

120 115 Lys Asp Thr Arg Leu Leu Thr Phe Lys Ala Phe Ala Gln Leu Phe Ile 135 140 Leu Gly Cys Ser Trp Val Leu Gly Ile Phe Gln Ile Gly Pro Val Ala 150 155 Gly Val Met Ala Tyr Leu Phe His His His Gln Gln Pro Ala Gly Gly 165 170 . 175 Leu His Leu Pro His Pro Leu Ser Ala Gln Arg Pro Gly Thr Arg Arg 185 Ile Gln Glu Val Asp His Trp Glu Asp Glu Ala Gln Leu Pro Val Pro 200 Asp Leu Lys Asp Leu Ala Val Leu His Ala Ile Arg Phe Gln Asp Gly 215 220 Leu Lys Ser Phe Leu Ala Phe Lys Tyr Ala Met Glu Pro Thr Val Gly 230 235 Gly Thr Ser Ser Phe Pro Cys Arg Glu Pro Tyr Pro * 250 252

<210> 1223 <211> 858 <212> PRT <213> Homo sapiens

<400> 1223 Met Lys Met Leu Thr Arg Leu Gln Val Leu Thr Leu Ala Leu Phe Ser 1 5 10 Lys Gly Phe Leu Leu Ser Leu Gly Asp His Asn Phe Leu Arg Arg Glu 25 Ile Lys Ile Glu Gly Asp Leu Val Leu Gly Gly Leu Phe Pro Ile Asn 40 Glu Lys Gly Thr Gly Thr Glu Glu Cys Gly Arg Ile Asn Glu Asp Arg 55 Gly Ile Gln Arg Leu Glu Ala Met Leu Phe Ala Ile Asp Glu Ile Asn 75 Lys Asp Asp Tyr Leu Leu Pro Gly Val Lys Leu Gly Val His Ile Leu Asp Thr Cys Ser Arg Asp Thr Tyr Ala Leu Glu Gln Ser Leu Glu Phe 105 Val Arg Ala Ser Leu Thr Lys Val Asp Glu Ala Glu Tyr Met Cys Pro 120 Asp Gly Ser Tyr Ala Ile Gln Glu Asn Ile Pro Leu Leu Ile Ala Gly 135 140 Val Ile Gly Gly Ser Tyr Ser Arg Val Ser Ile Gln Gly Ala Asn Leu 155 150 Leu Arg Leu Phe Gln Ile Pro Gln Ile Arg Tyr Ala Ser Thr Ser Ala 170 165 Lys Leu Ser Asp Lys Ser Arg Tyr Asp Tyr Phe Ala Arg Thr Val Pro 180 185 Pro Asp Phe Tyr Gln Ala Lys Ala Met Ala Glu Ile Leu Arg Phe Phe 200 Asn Trp Thr Tyr Val Ser Thr Val Ala Ser Glu Gly Asp Tyr Gly Glu 215 220 Thr Gly Ile Glu Ala Phe Glu Glu Glu Ala Arg Leu Arg Asn Ile Cys 235 230 Ile Ala Thr Ala Glu Lys Val Gly Arg Ser Asn Ile Arg Lys Ser Tyr 245 250

Asp	Ser	Val	Ile 260	Arg.	Glu	Leu	Leu	Gln 265	Lys	Pro	Asn	Ala	Arg 270	Val	Val
Val	Leu	Phe 275	Met	Arg	Ser	Asp	Asp 280	Ser	Arg	Glu	Leu	Ile 285	Ala	Ala	Ala
Ser			Asn	Ala	Ser			Trp	Val	Ala			Gly	Trp	${\tt Gl}_{Y}$
	290 Gln	Glu	Ser	Ile		295 Lys	Gly	Ser	Glu		300 Val	Ala	Tyr	Gly	Ala
305	_				310					315					320
Ile	Thr	Leu	Glu	Leu 325	Ala	Ser	Gln	Pro	Val 330	Arg	Gln	Phe	Asp	Arg 335	Tyr
Phe	Gln	Ser	Leu 340	Asn	Pro	Tyr	Asn	Asn 345	His	Arg	Asn	Pro	Trp 350	Phe	Arg
Asp	Phe	Trp 355	Glu	Gln	Lys	Phe	Gln 360	Cys	Ser	Leu	Gln	Asn 365	Lys	Arg	Asn
His	Ara		Val	Cvs	Asn	Lvs		Leu	Δla	Tle	Δsn		Ser	Δsn	ጥኒም
1110	370	9	val	Cyb	2105	375	*****	<u> </u>	7114		380	DCI	501	******	- y -
Glu		Glu	Ser	Lvs	Ile		Phe	Val	Val	Asn		Val	Tvr	Ala	Met
385				-	390					395			2 .		400
Ala	His	Ala	Leu	His	Lys	Met	Gln	Arg	Thr	Leu	Cys	Pro	Asn	Thr	Thr
				405	-			_	410		•			415	
Lys	Leu	Cys	Asp	Ala	Met	Lys	Ile	Leu	Asp	Gly	Lys	Lys	Leu	Tyr	Lys
			420					425					430		
Asp	Tyr	Leu 435	Leu	Lys	Ile	Asn	Phe 440	Thr	Ala	Pro	Phe	Asn 445	Pro	Asn	Lys
Asp	Ala	Asp	Ser	Ile	Val	Lys	Phe	Asp	Thr	Phe	Gly	Asp	Gly	Met	${ t Gly}$
	450					455					460				
Arg 465	Tyr	Asn	Val	Phe	Asn 470	Phe	Gln	Asn	Val	Gly 475	Gly	Lys	Tyr	Ser	Tyr 480
Leu	Lys	Val	Gly	His 485	Trp	Ala	Glu	Thr	Leu 490	Ser	Leu	Asp	Val	Asn 495	Ser
Ile	His	Trp	Ser 500	Arg	Asn	Ser	Val	Pro 505		Ser	Gln	Cys	Ser 510		Pro
Cys	Ala	Pro 515	Asn	Glu	Met	Lys	Asn 520		Gln	Pro	Gly	Asp 525	. – .	Cys	Cys
Trp	Ile		Ile	Pro	Cvs	Glu		Tvr	Glu	Tvr	Leu		Asp	Glu	Phe
	530	-2			-2-	535		- 1 -		-1-	540				
Thr	Cys	Met	Asp	Cys	Gly	Ser	Gly	Gln	Trp	Pro	Thr	Ala	Asp	Leu	Thr
545					550					555					560
Gly	Cys	Tyr	Asp	Leu 565	Pro	Glu	Asp	Tyr	Ile 570	Arg	Trp	Glu	Asp	Ala 575	Trp
Ala	Ile	Gly	Pro 580		Thr	Ile		Cys 585		Gly	Phe	Met	Cys 590	Thr	Cys
Met	Val	Val 595	Thr		Phe	Ile	Lys 600	His	Asn	Asn	Thr	Pro 605		Val	Lys
Ala	Ser		Arg	Glu	Leu	Cvs		Ile	Leu	Leu	Phe		Val	Glv	Len
	610	1	3			615	-1-				620	0-1		·]	
Ser		Cvs	Met	Thr	Phe		Phe	Ile	Ala	Lvs		Ser	Pro	Val	Ile
625	-	•			630					635					640
Cys	Ala	Leu	Arg	Arg	Leu	Gly	Leu	Gly	Ser	Ser	Phe	Ala	Ile	Cys	
				645		-		_	650					655	-
Ser	Ala	Leu	Leu 660	Thr	Lys	Thr	Asn	Cys 665	Ile	Ala	Arg	Ile	Phe 670	Asp	Gly
Val	Lys	Asn 675	Gly	Ala	Gln	Arg	Pro 680		Phe	Ile	Ser	Pro 685		Ser	Gln
Val	Phe		Cys	Leu	Glv	Leu		Leu	Val	Gln	Ile		Met	Val	Ser
	690		. 4		-4	695					700				
Val 705		Leu	Ile	Leu	Glu 710		Pro	Gly	Thr	Arg 715		Tyr	Thr	Leu	Ala 720
	Lys	Arq	Glu	Thr		Ile	Leu	Lys	Cys		Val	Lys	asA	Ser	
	-	_					-	-	-			-	-		

725 730 Met Leu Ile Ser Leu Thr Tyr Asp Val Ile Leu Val Ile Leu Cys Thr 740 745 Val Tyr Ala Phe Lys Thr Arg Lys Cys Pro Glu Asn Phe Asn Glu Ala 760 Lys Phe Ile Gly Phe Thr Met Tyr Thr Thr Cys Ile Ile Trp Leu Ala 775 780 Phe Leu Pro Ile Phe Tyr Val Thr Ser Ser Asp Tyr Arg Val Gln Thr 790 795 Thr Thr Met Cys Ile Ser Val Ser Leu Ser Gly Phe Val Val Leu Gly 805 810 Cys Leu Phe Ala Pro Lys Val His Ile Ile Leu Phe Gln Pro Gln Lys 820 825 Asn Val Val Thr His Arg Leu His Leu Asn Arg Phe Ser Val Ser Gly 835 840 Thr Gly Thr His Ile Leu Ser Val Leu * 855 857

<210> 1224

<211> 69

<212> PRT

<213> Homo sapiens

<400> 1224

 Met
 Ser
 His
 Met
 Val
 Pro
 Leu
 Ala
 Leu
 Leu
 Leu
 Leu
 Pro
 Thr
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 Thr
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 Thr
 15
 Thr
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 Thr
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 15
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 16
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 Thr
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<210> 1225

<211> 55

<212> PRT

<213> Homo sapiens

<400> 1225

 Met Cys
 Tyr His Thr Trp Leu Ile Phe Ile Phe Leu Val Glu Met Gly

 1
 5
 10
 15

 Phe Tyr His Val Gly Gln Ala Gly Phe Lys Leu Leu Ala Ser Ser Gly
 20
 25
 30

 Pro Pro Ala Ser Ala Ser Gln Ser Ala Gly Ile Thr Gly Val Ser His
 35
 40
 45

 His Ala Arg Pro Thr Phe
 *

<210> 1226

<211> 51 <212> PRT <213> Homo sapiens

<400> 1226

<210> 1227 <211> 47 <212> PRT <213> Homo sapiens

<210> 1228 <211> 60 <212> PRT <213> Homo sapiens

<210> 1229 <211> 52 <212> PRT <213> Homo sapiens

<400> 1229
Met Cys Glu Ser Thr Glu Leu Asn Met Thr Phe His Leu Phe Ile Val

<210> 1230 <211> 362 <212> PRT <213> Homo sapiens

<400> 1230 Met Pro Val Ile Trp Ser Ala Leu Ser Ala Val Leu Leu Leu Ala Ser 10 Ser Tyr Phe Val Gly Ala Leu Ile Val His Ala Asp Cys Phe Leu Met 25 Arg Asn His Thr Ile Thr Glu Gln Pro Met Cys Phe Gln Arg Thr Thr 40 Pro Leu Ile Leu Gln Glu Val Ala Ser Phe Leu Lys Arg Asn Lys His 55 Gly Pro Phe Leu Leu Phe Val Ser Phe Leu His Val His Ile Pro Leu 70 75 Ile Thr Met Glu Asn Phe Leu Gly Lys Ser Leu His Gly Leu Tyr Gly 90 85 Asp Asn Val Lys Glu Met Asp Trp Met Val Gly Arg Ile Leu Asp Thr 100 105 Leu Asp Val Glu Gly Leu Ser Asn Ser Thr Leu Ile Tyr Phe Thr Ser 120 125 Asp His Gly Gly Ser Leu Glu Asn Gln Leu Gly Asn Thr Gln Tyr Gly 135 140 Gly Trp Asn Gly Ile Tyr Lys Gly Gly Lys Gly Met Gly Gly Trp Glu 150 155 Gly Gly Ile Arg Val Pro Gly Ile Phe Arg Trp Pro Gly Val Leu Pro 170 Ala Gly Arg Val Ile Gly Glu Pro Thr Ser Leu Met Asp Val Phe Pro 185 Thr Val Val Arg Leu Ala Gly Ser Glu Val Pro Gln Asp Arg Val Ile 200 Asp Gly Gln Asp Leu Leu Pro Leu Leu Gly Thr Ala Gln His Ser 215 Asp His Glu Phe Leu Met His Tyr Cys Glu Arg Phe Leu His Ala Ala 230 235 Arg Trp His Gln Arg Asp Arg Gly Thr Met Trp Lys Val His Phe Val 245 250 Thr Pro Val Phe Gln Pro Arg Gly Ser Arg Cys Leu Leu Trp Lys Glu 260 265 Lys Val Cys Pro Cys Phe Gly Glu Lys Ser Ser Pro Pro Arg Ser His 285 280 Pro Cys Phe Phe Asp Leu Ser Arg Ala Pro Ser Glu Thr His Ile Leu 295 300 Thr Pro Ala Ser Glu Pro Val Phe Tyr Gln Val Met Glu Arg Ser Pro 310 315 Ala Gly Gly Val Gly Thr Pro Ala Asp Thr Gln Pro Ser Ser Ala 325 330

<210> 1231 <211> 53 <212> PRT <213> Homo sapiens

<400> 1231

 Met
 Leu
 Arg
 Leu
 Gly
 Val
 Ala
 Phe
 His
 Met
 Glu
 Leu
 Leu
 Cys
 Arg
 Gly

 Arg
 Leu
 Leu
 Leu
 Ile
 Pro
 Thr
 Ala
 Glu
 Thr
 Arg
 Cys
 Asp
 His
 Arg

 Arg
 Leu
 Leu
 Leu
 Ser
 Asn
 Thr
 Leu
 Asp
 Lys
 His

 35
 40
 45
 45

 Gln
 Glu
 Pro
 His
 *

 50
 52

<210> 1232 <211> 56 <212> PRT <213> Homo sapiens

<210> 1233 <211> 56 <212> PRT <213> Homo sapiens

<210> 1234 <211> 125 <212> PRT <213> Homo sapiens

<210> 1235 <211> 72 <212> PRT <213> Homo sapiens

<210> 1236 <211> 48 <212> PRT <213> Homo sapiens

Arg Ala Gly Gly Leu Gly Phe Thr His Cys Gln Ala Asn Ser Thr Thr 35 40 45 48

<210> 1237 <211> 208 <212> PRT <213> Homo sapiens

<400> 1237 Met Ala Phe Leu Arg Lys Val Tyr Ser Ile Leu Ser Leu Gln Val Leu Leu Thr Thr Val Thr Ser Thr Val Phe Leu Tyr Phe Glu Ser Val Arg 20 25 Thr Phe Val His Glu Ser Pro Ala Leu Ile Leu Leu Phe Ala Leu Gly 40 Ser Leu Gly Leu Ile Phe Ala Leu Ile Leu Asn Arg His Lys Tyr Pro 55 Leu Asn Leu Tyr Leu Leu Phe Gly Phe Thr Leu Leu Glu Ala Leu Thr 75 70 Val Ala Val Val Thr Phe Tyr Asp Val Tyr Ile Ile Leu Gln Ala 90 85 Phe Ile Leu Thr Thr Thr Val Phe Phe Gly Leu Thr Val Tyr Thr Leu 100 105 Gln Ser Lys Lys Asp Phe Ser Lys Phe Gly Ala Gly Leu Phe Ala Leu 120 125 Leu Trp Ile Leu Cys Leu Ser Gly Phe Leu Lys Phe Phe Phe Tyr Ser 140 135 Glu Ile Met Glu Leu Val Leu Ala Ala Gly Ala Leu Leu Phe Cys 155 150 Gly Phe Ile Ile Tyr Asp Thr His Ser Leu Met His Lys Leu Ser Pro 170 165 Glu Glu Tyr Val Leu Ala Ala Ile Ser Leu Tyr Leu Asp Ile Ile Asn 185 Leu Phe Leu His Leu Leu Arg Phe Leu Glu Ala Val Asn Lys Lys * 205 200 195

<210> 1238 <211> 173 <212> PRT <213> Homo sapiens

70 75 Asn Phe Gly Phe Ser Leu Leu Arg Lys Ile Ser Met Arg His Asp Gly 90 85 Asn Met Val Phe Ser Pro Phe Gly Met Ser Leu Ala Met Thr Gly Leu 100 105 Met Leu Gly Ala Thr Gly Pro Thr Glu Thr Gln Ile Lys Arg Gly Leu 120 His Leu Gln Ala Leu Lys Pro Thr Lys Pro Gly Leu Leu Pro Ser Leu 135 140 Phe Lys Gly Leu Arg Glu Thr Leu Ser Arg Asn Leu Glu Leu Gly Leu 150 155 Thr Ala Gly Glu Phe Cys Leu His Pro Gln Gly Phe *

<210> 1239 <211> 357 <212> PRT <213> Homo sapiens

<400> 1239 Met Ala Phe Leu Gly Leu Phe Ser Leu Leu Val Leu Gln Ser Met Ala 1 5 10 15 Thr Gly Ala Thr Phe Pro Glu Glu Ala Ile Ala Asp Leu Ser Val Asn 25 Met Tyr Asn Arg Leu Arg Ala Thr Gly Glu Asp Glu Asn Ile Leu Phe 40 Ser Pro Leu Ser Ile Ala Leu Ala Met Gly Met Met Glu Leu Gly Ala 55 Gln Gly Ser Thr Gln Lys Glu Ile Arg His Ser Met Gly Tyr Asp Ser 70 75 Leu Lys Asn Gly Glu Glu Phe Ser Phe Leu Lys Glu Phe Ser Asn Met 90 85 Val Thr Ala Lys Glu Ser Gln Tyr Val Met Lys Ile Ala Asn Ser Leu 100 105 Phe Val Gln Asn Gly Phe His Val Asn Glu Glu Phe Leu Gln Met Met 120 Lys Lys Tyr Phe Asn Ala Ala Val Asn His Val Asp Phe Ser Gln Asn 135 140 Val Ala Val Ala Asn Tyr Ile Asn Lys Trp Val Glu Asn Asn Thr Asn 150 155 160 Asn Leu Val Lys Asp Leu Val Ser Pro Arg Asp Phe Asp Ala Ala Thr 170 Tyr Leu Ala Leu Ile Asn Ala Val Tyr Phe Lys Gly Asn Trp Lys Ser 185 Gln Phe Arg Pro Glu Asn Thr Arg Thr Phe Ser Phe Thr Lys Asp Asp 200 Glu Ser Glu Val Gln Ile Pro Met Met Tyr Gln Gln Gly Glu Phe Tyr 215 Tyr Gly Glu Phe Ser Asp Gly Ser Asn Glu Ala Gly Gly Ile Tyr Gln 230 235 Val Leu Glu Ile Pro Tyr Glu Gly Asp Glu Ile Ser Met Met Leu Val 250 245 Leu Ser Arg Gln Glu Val Pro Leu Ala Thr Leu Glu Pro Leu Val Lys 265 Ala Gln Leu Val Glu Glu Trp Ala Asn Ser Val Lys Lys Gln Lys Val 275 280 285

<210> 1240 <211> 707 <212> PRT <213> Homo sapiens

<400> 1240 Met Leu Ser Leu Arg Arg Cys Thr Ser Met Arg Leu Cys Leu Ser Ser 10 Ser Leu Ala Ser Pro Cys Ser Thr Met Leu Ser Thr Val Val Leu Tyr 25 Lys Val Cys Asn Ser Phe Val Glu Met Gly Ser Ala Asn Val Gln Ala 40 Thr Asp Tyr Leu Lys Gly Val Ala Ser Leu Phe Val Val Ser Leu Gly 55 Gly Ala Ala Val Gly Leu Val Phe Ala Phe Leu Leu Ala Leu Thr Thr 75 Arg Phe Thr Lys Arg Val Arg Ile Ile Glu Pro Leu Leu Val Phe Leu 85 90 Leu Ala Tyr Ala Ala Tyr Leu Thr Ala Glu Met Ala Ser Leu Ser Ala 105 Ile Leu Ala Val Thr Met Cys Gly Leu Gly Cys Lys Lys Tyr Val Glu 120 Ala Asn Ile Ser His Lys Ser Arg Thr Thr Val Lys Tyr Thr Met Lys 135 140 Thr Leu Ala Ser Cys Ala Glu Thr Val Ile Phe Met Leu Leu Gly Ile 150 155 Ser Thr Val Asp Ser Ser Lys Trp Ala Trp Asp Ser Gly Leu Val Leu 165 170 Gly Thr Leu Ile Phe Ile Leu Phe Phe Arg Ala Leu Gly Val Val Leu 185 180 Gln Thr Trp Val Leu Asn Gln Phe Arg Leu Val Pro Leu Asp Lys Ile 200 Asp Gln Val Val Met Ser Tyr Gly Gly Leu Arg Gly Ala Val Ala Phe 215 220 Ala Leu Val Ile Leu Leu Asp Arg Thr Lys Val Pro Ala Lys Asp Tyr 230 235 Phe Val Ala Thr Thr Ile Val Val Phe Phe Thr Val Ile Val Gln 245 250 Gly Leu Thr Ile Lys Pro Leu Val Lys Trp Leu Lys Val Lys Arg Ser 265 270 Glu His His Lys Pro Thr Leu Asn Gln Glu Leu His Glu His Thr Phe 280 Asp His Ile Leu Ala Ala Val Glu Asp Val Val Gly His His Gly Tyr 295 His Tyr Trp Arg Asp Arg Trp Glu Gln Phe Asp Lys Lys Tyr Leu Ser

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305					310					315					320
Gln	Leu	Leu	Met	Arg	Arg	Ser	Ala	Tyr	Arg	Ile	Arg	Asp	${\tt Gln}$	Ile	\mathtt{Trp}
				325					330					335	
Asp	Val	Tyr	Tyr	Arg	Leu	Asn	Ile		Asp	Ala	Ile	Ser		Val	Asp
~ 3	~ 1		340	7	.	0	0	345	~1	*		T	350	0	36- 1
GIN	GTĀ	355	His	vaı	Leu	ser	360	Thr	GIĀ	ьeu	Thr	ьеи 365	Pro	ser	Met
Dro	Sor		Asn	Car	Va l	ת דות		Thr	Sar	TeV	Thr		T.011	T.211	7/ 2/ (1
PIO	370	Arg	ASII	Ser	vaı	375	GIU	1111	361	vai	380	WOII	цец	пец	Arg
Glu		Glv	Ser	Glv	Ala		Leu	Asp	Leu	Gln	_	Ile	asp	Thr	Val
385		,			390			1		395					400
Arg	Ser	Gly	Arg	Asp	Arg	Glu	Asp	Ala	Val	Met	His	His	Leu	Leu	Cys
				405					410					415	
Gly	Gly	Leu	Tyr	Lys	Pro	Arg	Arg		Tyr	Lys	Ala	Ser	Cys	Ser	Arg
•	_,		420		_		~ 7	425	_	~-3	_	_	430		
His	Phe		Ser	GLu	Asp	Ala		GLu	Arg	GIn	Asp		Glu	Val	Phe
Gln	Gln	435	Met	Tare	Λrσ	Λνα	440	Glu	Sar	Dha	Tare	445	ሞኩዮ	Lve	Иic
GIII	450	ASII	Mec	шуэ	Arg	455	nea	GIU	Jer	FIIC	460	Der	1111	цуз	1113
Asn		Cvs	Phe	Thr	Lvs		Lvs	Pro	Arq	Pro		Lvs	Thr	Glv	Arq
465		4			470		4			475	_	_		-	480
Arg	Lys	Lys	Asp	Gly	Val	Ala	Asn	Ala	Glu	Ala	Thr	Asn	Gly	Lys	His
				485					490					495	
Arg	Gly	Leu	Gly	Phe	Gln	Asp	Thr		Ala	Val	Ile	Leu		Val	Glu
_	~1	~7	500	~7	~ 3	~ 3	~	505	_	~	~1		510	T	~1
Ser	GIU	515	Glu	GIU	Glu	GIU	520	Asp	ser	ser	GIU	525	GIU	гуѕ	GIU
Δαρ	Δen		Gly	Tle	Tle	Dhe		Δla	Δra	Δla	Thr		Glu	Val	T.em
пор	530	014	C±y			535	VUL	1114	9	1114	540	501	<u> </u>	Vul	LCu
Gln		Gly	Lys	Val	Ser	Gly	Ser	Leu	Glu	Val	Cys	Pro	Ser	Pro	Arg
545		•	_		550	_				555	_				560
Ile	Ile	Pro	Pro	Ser	Pro	Thr	Cys	Ala	Glu	Lys	Glu	Leu	Pro	Trp	Lys
_	.			565	_				570					575	
Ser	GLY	Gln	Gly	Asp	Leu	Ala	Val		Val	Ser	Ser	GLu		Thr	Lys
Tla	7727	Dro	580 Val	7) an	Mot	Gl n	Thr	585	Trn	λen	aln	Sar	590	Car	Car
116	vaı	595	vai	ASP	Met	GIII	600	GTA	пр	ASII	GIII	605	116	261	Ser
Leu	Glu		Leu	Ala	Ser	Pro		Cys	Asn	Gln	Ala		Ile	Leu	Thr
	610					615		-			620				
Cys	Leu	Pro	Pro	His	Pro	Arg	Gly	Thr	Glu	Glu	Pro	Gln	Val	Pro	Leu
625					630					635					640
His	Leu	Pro	Ser		Pro	Arg	Ser	Ser		Ala	Phe	Pro	Pro		Leu
77-	T	71 -	~1	645	a	*	0	~1	650	0	77-	71	T	655	71
нта	ьуѕ	АТА	660	arg	ser	arg	ser	665	ser	ser	АТА	Asp	ьеи 670	Pro	GIN
Gln	Gln	Glu	Leu	Gln	Pro	T.e.v	Met		Hic	Tave	Aso	Hie		His	T,@11
0111	U111	675	Leu	U-111	110		680	CT Y	****	دىرى	LISP	685	****		u
Ser	Pro		Thr	Ala	Thr	Ser		Trp	Cys	Ile	Gln		Asn	Arg	Gly
	690	•				695		-	-		700			_	•
Ser	Arg	Leu													
705		707													

<210> 1241

<211> 98 <212> PRT

<213> Homo sapiens

<400> 1241 Met Ala Phe Arg Thr Phe Ser Trp Ile Phe Ser Gly Leu Leu Ser Pro Thr Leu Ala Ser Pro Ser Val Ser Met Met Thr Met Glu Val Leu Leu 25 Ser Gly Ile Leu Cys Ser Ser Arg Ala Leu Phe Ser Ile Leu Met Pro 40 4.5 Leu Ser Ser Pro Ser Leu Met Leu Val Ile Pro Leu Ser Ser Met Leu 5.5 Phe Thr Asn Val Leu Ala Ser Trp Arg Phe Ser Gly Val Ala Trp Thr 70 75 Lys Cys Ser Phe His Val Asp Thr Ser Pro Leu Asn Arg Met Lys Phe 85 90 Arg * 97

<210> 1242 <211> 422 <212> PRT <213> Homo sapiens

<400> 1242 Met Val Leu Trp Glu Ser Pro Arg Gln Cys Ser Ser Trp Thr Leu Cys 10 Glu Gly Phe Cys Trp Leu Leu Leu Pro Val Met Leu Leu Ile Val 20 2.5 Ala Arg Pro Val Lys Leu Ala Ala Phe Pro Thr Ser Leu Ser Asp Cys 40 Gln Thr Pro Thr Gly Trp Asn Cys Ser Gly Tyr Asp Asp Arg Glu Asn 55 Asp Leu Phe Leu Cys Asp Thr Asn Thr Cys Lys Phe Asp Gly Glu Cys 70 75 Leu Arg Ile Gly Asp Thr Val Thr Cys Val Cys Gln Phe Lys Cys Asn 85 90 Asn Asp Tyr Val Pro Val Cys Gly Ser Asn Gly Glu Ser Tyr Gln Asn 100 105 110 Glu Cys Tyr Leu Arg Gln Ala Ala Cys Lys Gln Gln Ser Glu Ile Leu 120 Val Val Ser Glu Gly Ser Cys Ala Thr Asp Ala Gly Ser Gly Ser Gly 135 Asp Gly Val His Glu Gly Ser Gly Glu Thr Ser Gln Lys Glu Thr Ser 150 155 Thr Cys Asp Ile Cys Gln Phe Gly Ala Glu Cys Asp Glu Asn Ala Glu 170 Asp Val Trp Cys Val Cys Asn Ile Asp Cys Ser Gln Thr Asn Phe Asn 185 Pro Leu Cys Ala Ser Asp Gly Lys Ser Tyr Asp Asn Ala Cys Gln Ile 200 205 Lys Glu Ala Ser Cys Gln Lys Gln Glu Lys Ile Glu Val Leu Ser Leu 215 220 Gly Arg Cys Gln Asp Asn Thr Thr Thr Thr Thr Lys Ser Glu Asp Gly 230 235 His Tyr Ala Arg Thr Asp Tyr Ala Glu Asn Ala Asn Lys Leu Glu Glu 245 250 Ser Ala Arg Glu His His Ile Pro Cys Pro Glu His Tyr Asn Gly Phe

265 260 Cys Met His Gly Lys Cys Glu His Ser Ile Asn Met Gln Glu Pro Ser 280 Cys Arg Cys Asp Ala Gly Tyr Thr Gly Gln His Cys Glu Lys Lys Asp 295 300 Tyr Ser Val Leu Tyr Val Val Pro Gly Pro Val Arg Phe Pro Val Cys 310 315 Leu Asn Arg Ser Cys Asp Trp Asn Asn Ser Asp Cys Cys His Leu Cys 325 330 Gly Gly Pro Leu His His Lys Glu Met Pro Pro Glu Ala Asn Arg Ile 345 340 Pro Pro Asp Arg Ser Lys Ile Pro Gly His Tyr Ser Ser Arg Gln Tyr 360 Asn Lys Ser Arg Pro Thr Arg Leu Ile Leu Lys Gly Ala Cys Phe His 370 375 Ser Gly Trp Thr Thr Glu Ser Leu Asp Tyr Thr Ile Gln Tyr Tyr Arg 390 395 Gln Lys Asn Lys Thr Arg Asp Leu Thr His Val Cys Leu Ala Phe Val 405 410 Gly Asn Leu His Gln * 420 421

<210> 1243 <211> 46

<212> PRT

<213> Homo sapiens

<400> 1243

<210> 1244

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1244

 Met Val Leu Ser Ala Pro Ser Leu Trp Pro Cys Ser Ser Phe Ser Ile

 1
 5
 10
 15

 Ser Cys Leu His Val Gly Leu Thr Ala Phe Leu Phe Gln Val Ala Phe
 20
 25
 30

 Leu Cys Leu Leu Cys Cys Val Glu Leu Leu Leu Asp Val
 *

 35
 40
 45

<210> 1245

<211> 244

<212> PRT

<213> Homo sapiens

<400> 1245 Met Ala Gly Val Ile Ala Gly Leu Leu Met Phe Ile Ile Ile Leu Leu Gly Val Met Leu Thr Ile Lys Arg Arg Arg Asn Ala Tyr Ser Tyr Ser 2.0 25 Tyr Tyr Leu Lys Leu Ala Lys Lys Gln Lys Glu Thr Gln Ser Gly Ala 40 Gln Arg Glu Met Gly Pro Val Ala Ser Ala Asp Lys Pro Thr Thr Lys 55 60 Leu Ser Ala Ser Arg Asn Asp Glu Gly Phe Ser Ser Ser Ser Gln Asp 70 75 Val Asn Gly Phe Asn Gly Ser Arg Gly Glu Leu Ser Gln Pro Thr Leu 85 90 Thr Ile Gln Thr His Pro Tyr Arg Thr Cys Asp Pro Val Glu Met Ser 105 Tyr Pro Arg Asp Gln Phe Gln Pro Ala Ile Arg Val Ala Asp Leu Leu 120 Gln His Ile Thr Gln Met Lys Arg Gly Gln Gly Tyr Gly Phe Lys Glu 135 Glu Tyr Glu Ala Leu Pro Glu Gly Gln Thr Ala Ser Trp Asp Thr Ala 150 155 Lys Glu Asp Glu Asn Arg Asn Lys Asn Arg Tyr Gly Asn Ile Ile Ser 165 170 Tyr Asp His Ser Arg Val Arg Leu Leu Val Leu Asp Gly Asp Pro His 185 Ser Asp Tyr Ile Asn Ala Asn Tyr Ile Asp Gly Tyr His Arg Pro Arg 200 205 His Tyr Ile Ala Thr Gln Gly Pro Met Gln Glu Thr Val Lys Asp Phe 215 220 Trp Arg Met Ile Trp Gln Glu Asn Ser Ala Ser Ile Val Met Val Thr 230 235 Asn Pro Gly * 243

<210> 1246 <211> 565 <212> PRT

<213> Homo sapiens

<400> 1246 Met Ala Val Phe Arg Ser Gly Leu Leu Val Leu Thr Thr Pro Leu Ala 10 Ser Leu Ala Pro Arg Leu Ala Ser Ile Leu Thr Ser Ala Ala Arg Leu 25 Val Asn His Thr Leu Tyr Val His Leu Gln Pro Gly Met Ser Leu Glu 40 Gly Pro Ala Gln Pro Gln Tyr Ser Pro Val Gln Ala Thr Phe Glu Val 55 Leu Asp Phe Ile Thr His Leu Tyr Ala Gly Ala Asp Val His Arg His 75 70 Leu Asp Val Arg Ile Leu Leu Thr Asn Ile Arg Thr Lys Ser Thr Phe 85 90 Leu Pro Pro Leu Pro Thr Ser Val Gln Asn Leu Ala His Pro Pro Glu

			7.00					105					110		
Val	Val	T.e.11	100 Thr	Asp	Phe	Gln	Thr	105 Leu	Asp	Glv	Ser	Gln	110 Tvr	Asn	Pro
• • • •	· ~ _	115				0	120			1		125	- 1 -		
Val	Lys 130	Gln	Gln	Leu	Val	Arg 135	Tyr	Ala	Thr	Ser	Cys 140	Tyr	Ser	Cys	Cys
Pro 145	Arg	Leu	Ala	Ser	Val 150	Leu	Leu	Tyr	Ser	Asp 155	Tyr	Gly	Ile	Gly	Glu 160
			Glu	165					170					175	
			Val 180					185					190		
_		195	Val				200					205			
	210		Ser			215					220				
G1y 225	Val	Ala	Asp	Lys	230	Leu	Leu	Lys	Ser	ட்ys 235	ьeu	Leu	Pro	GIU	ьеи 240
			Tyr	245					250					255	
			Pro 260					265					270		
_	_	275	Ala				280					285			
	290		Tyr			295					300				
Asn 305	Asp	Leu	Glu	Glu	Leu 310	Ala	Leu	Tyr	Gln	Ile 315	Gln	Leu	Leu	Lys	Asp 320
Leu	Arg	His	Thr	Glu 325	Asn	Glu	Glu	Asp	130 130	Val	Ser	Ser	Ser	Ser 335	Phe
_		_	Met 340		_			345					350		
		355	Thr	-		_	360		-			365			
	370	_	Ser			375					380				
385			Ser		390					395					400
		-	Gln	405					410	_		_		415	
_	_	_	11e 420			_	_	425					430		
	_	435	Gln				440					445			
	450		Ala			455					460				
465		_	Val		470	•				475					480
			His	485					490					495	
	_	_	Ile 500			_	_	505					510		
		515	Gln				520					525			
	530		Leu			535					540				
Val 545	Glu	Lys	Ala	Trp	Ala 550	Leu	Leu	Gln	Lys	Arg 555	Ile	Pro	Lys	Thr	His 560
Gln	Ala	Leu	Asp 564	*											

<210> 1247 <211> 737 <212> PRT <213> Homo sapiens

<400> 1247 Met Phe Pro Ala Gly Pro Pro Trp Pro Arg Val Arg Val Val Gln Val Leu Trp Ala Leu Leu Ala Val Leu Leu Ala Ser Trp Arg Leu Trp Ala 25 Ile Lys Asp Phe Gln Glu Cys Thr Trp Gln Val Val Leu Asn Glu Phe 40 Lys Arg Val Gly Glu Ser Gly Val Ser Asp Ser Phe Phe Glu Glu Glu 55 Pro Val Asp Thr Val Ser Ser Leu Phe His Met Leu Val Asp Ser Pro 75 70 Ile Asp Pro Ser Glu Lys Tyr Leu Gly Phe Pro Tyr Tyr Leu Lys Ile 85 90 Asn Tyr Ser Cys Glu Glu Lys Pro Ser Glu Asp Leu Val Arg Met Gly 100 105 His Leu Thr Gly Leu Lys Pro Leu Val Leu Val Thr Phe Gln Ser Pro 120 Val Asn Phe Tyr Arg Trp Lys Ile Glu Gln Leu Gln Ile Gln Met Glu 135 Ala Ala Pro Phe Arg Ser Lys Gly Gly Pro Gly Gly Gly Arg Asp 150 155 Arg Asn Leu Ala Gly Met Asn Ile Asn Gly Phe Leu Lys Arg Asp Arg 165 170 Asp Asn Asn Ile Gln Phe Thr Val Gly Glu Leu Phe Asn Leu Met 185 Pro Gln Tyr Phe Val Gly Val Ser Ser Arg Pro Leu Trp His Thr Val 200 Asp Gln Ser Pro Val Leu Ile Leu Gly Gly Ile Pro Asn Glu Lys Tyr 215 Val Leu Met Thr Asp Thr Ser Phe Lys Asp Phe Ser Leu Val Glu Val 230 235 Asn Gly Val Gly Gln Met Leu Ser Ile Asp Ser Cys Trp Val Gly Ser 245 250 Phe Tyr Cys Pro His Ser Gly Phe Thr Ala Thr Ile Tyr Asp Thr Ile 265 Ala Thr Glu Ser Thr Leu Phe Ile Arg Gln Asn Gln Leu Val Tyr Tyr 280 Phe Thr Gly Thr Tyr Thr Leu Tyr Glu Arg Asn Arg Gly Ser Gly 295 300 Glu Cys Ala Val Ala Gly Pro Thr Pro Gly Glu Gly Thr Leu Val Asn 315 310 Pro Ser Thr Glu Gly Ser Trp Ile Arg Val Leu Ala Ser Glu Cys Ile 325 330 Lys Lys Leu Cys Pro Val Tyr Phe His Ser Asn Gly Ser Glu Tyr Ile 345 Met Ala Leu Thr Thr Gly Lys His Glu Gly Tyr Val His Phe Gly Thr 360 Ile Arg Val Thr Thr Cys Ser Ile Ile Trp Ser Glu Tyr Ile Ala Gly 375 380 Glu Tyr Thr Leu Leu Leu Val Glu Ser Gly Tyr Gly Asn Ala Ser

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390
                                  395
Lys Arg Phe Gln Val Val Ser Tyr Asn Thr Ala Ser Asp Asp Leu Glu
             405
                              410
Leu Leu Tyr His Ile Pro Glu Phe Ile Pro Glu Ala Arg Gly Leu Glu
          420
                        425
Phe Leu Met Ile Leu Gly Thr Glu Ser Tyr Thr Ser Thr Ala Met Ala
                        440
                                         445
Pro Lys Gly Ile Phe Cys Asn Pro Tyr Asn Asn Leu Ile Phe Ile Trp
          455
Gly Asn Phe Leu Leu Gln Ser Ser Asn Lys Glu Asn Phe Ile Tyr Leu
465 470
                                 475
Ala Asp Phe Pro Lys Glu Leu Ser Ile Lys Tyr Met Ala Arg Ser Phe
              485
                               490
Arg Gly Ala Val Ala Ile Val Thr Glu Thr Glu Glu Ile Trp Tyr Leu
                            505
Leu Glu Gly Ser Tyr Arg Val Tyr Gln Leu Phe Pro Ser Lys Gly Trp
                        520
Gln Val His Ile Ser Leu Lys Leu Met Gln Gln Ser Ser Leu Tyr Ala
                  535
Ser Asn Glu Thr Met Leu Thr Leu Phe Tyr Glu Asp Ser Lys Leu Tyr
                550
                                  555
Gln Leu Val Tyr Leu Met Asn Asn Gln Lys Gly Gln Leu Val Lys Arg
             565 570
Leu Val Pro Val Glu Gln Leu Leu Met Tyr Gln Gln His Thr Ser His
                     585
          580
Tyr Asp Leu Glu Arg Lys Gly Gly Tyr Leu Met Leu Ser Phe Ile Asp
                        600
Phe Cys Pro Phe Ser Val Met Arg Leu Arg Ser Leu Pro Ser Pro Gln
                    615
                                  620
Arg Tyr Thr Arg Gln Glu Arg Tyr Arg Ala Arg Pro Pro Arg Val Leu
                                635 640
                630
Glu Arg Ser Gly Phe Pro Gln Gly Glu Leu Ala Arg His Leu Pro Gly
                              650
             645
Pro Gly Leu Pro Ala Val Ala Leu Arg Val Arg Gln Ala Val
                          665
Arg Gly Pro Gly Ala Arg Pro His Leu Ala Leu Val Gly Glu Gln Gln
              680
                                         685
Thr Arg Pro Gly Leu Leu Leu Leu Gly Glu Gln Leu Ala Lys Arg
              695
                                      700
Gly Arg Arg Val His Arg Asn Gly Gln Leu Arg Lys Asp Leu Gln Pro
                                715
                710
Arg Val Arg Val Arg Ala Ala Gly Ala His Phe Pro Gly Gln Gly His
                               730
                                                 735 736
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<210> 1248 <211> 175 <212> PRT <213> Homo sapiens

 $<\!400\!>$ 1248
Met Gly Trp Val Trp Thr Leu Cys Thr Ala Ser Ala Cys Leu Thr Leu 1 5 10 15
Leu Phe Trp Ser Gln Thr Pro Gly Lys Ala Phe Gln Ile Pro Cys Pro 20 30

Pro Pro His Leu Ser His Trp Cys Leu Ser Pro Met Gln Met Asp Asp Gly Cys Ala Arq Leu Cys Val Leu Trp Thr Ala Trp Met Arg Trp Arg Val Leu Met Cys Ser Cys Arg Val Trp Ala Thr Asp Leu Gly Ile Phe 70 75 Leu Gly Val Ala Leu Gly Asn Glu Pro Leu Glu Met Trp Pro Leu Thr 85 90 Gln Asn Glu Glu Cys Thr Val Thr Gly Phe Leu Arg Asp Lys Leu Gln 105 100 Tyr Arg Ser Arg Leu Gln Tyr Met Lys His Tyr Phe Pro Ile Asn Tyr 120 125 Lys Ile Arg Val Pro Tyr Glu Gly Val Phe Arg Ile Ala Asn Val Thr 135 140 Arg Leu Arg Ala Gln Gly Ser Glu Arg Glu Leu Arg Tyr Leu Gly Val 150 155 Leu Val Ser Leu Ser Ala Thr Glu Ser Val His Asp Glu Leu Leu 170

<210> 1249 <211> 68 <212> PRT

<213> Homo sapiens

<210> 1250 <211> 209 <212> PRT

<213> Homo sapiens

Leu Leu Lys Asp Glu Leu Leu Met Pro Ser Val Val Thr Thr Met

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90 85 Ala Phe Phe Ile Ala Cys Val Thr Ser Phe Ser Ile Phe Glu Lys Thr 105 100 Ser Glu Glu Glu Leu Gln Leu Lys Ser Phe Ser Ile Ser Val Arg Lys 115 120 Tyr Leu Pro Cys Phe Thr Phe Leu Ser Arg Ile Ile Gln Tyr Leu Phe 135 140 Leu Ile Ser Val Ile Thr Met Val Leu Leu Thr Leu Met Thr Val Thr 150 155 Leu Asp Pro Pro Gln Lys Leu Pro Asp Leu Phe Ser Val Leu Val Cys 170 Phe Val Ser Cys Leu Asn Phe Leu Phe Phe Leu Val Tyr Phe Asn Ile 185 Ile Ile Met Trp Asp Ser Lys Ser Gly Arg Asn Gln Lys Lys Ile Ser 200 205

<210> 1251 <211> 58 <212> PRT <213> Homo sapiens

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<213> Homo sapiens

> <210> 1254 <211> 209 <212> PRT <213> Homo sapiens

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<210> 1255 <211> 730 <212> PRT <213> Homo sapiens

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Ser Glu Tyr Thr Ser Leu Ile Pro Asn Leu Arg Asn Val Val Ala Leu
           420
                        425
Asp Thr Glu Val Ala Ser Asn Arg Ile Tyr Trp Ser Asp Leu Ser Gln
                           440
Arg Met Ile Cys Ser Thr Gln Leu Asp Arg Ala His Gly Val Ser Ser
                       455
                                           460
Tyr Asp Thr Val Ile Ser Arg Asp Ile Gln Ala Pro Asp Gly Leu Ala
                   470
                                       475
Val Asp Trp Ile His Ser Asn Ile Tyr Trp Thr Asp Ser Val Leu Gly
               485
                                   490
Thr Val Ser Val Ala Asp Thr Lys Gly Val Lys Arg Lys Thr Leu Phe
           500
                               505
Arg Glu Asn Gly Ser Lys Pro Arg Ala Ile Val Val Asp Pro Val His
       515
                           520
                                               525
Gly Phe Met Tyr Trp Thr Asp Trp Gly Thr Pro Ala Lys Ile Lys Lys
                       535
                                           540
Gly Gly Leu Asn Gly Val Asp Ile Tyr Ser Leu Val Thr Glu Asn Ile
                   550
                                       555
Gln Trp Pro Asn Gly Ile Thr Leu Asp Leu Leu Ser Gly Arg Leu Tyr
               565
                                   570
Trp Val Asp Ser Lys Leu His Ser Ile Ser Ser Ile Asp Val Asn Gly
                               585
Gly Asn Arg Lys Thr Ile Leu Glu Asp Glu Lys Arg Leu Ala His Pro
                           600
Phe Ser Leu Ala Val Phe Glu Asp Lys Val Phe Trp Thr Asp Ile Ile
                       615
                                           620
Asn Glu Ala Ile Phe Ser Ala Asn Arg Leu Thr Gly Ser Asp Val Asn
                   630
                                       635
Leu Leu Ala Glu Asn Leu Leu Ser Pro Glu Asp Met Val Leu Phe His
               645
                                   650
Asn Leu Thr Gln Pro Arg Gly Val Asn Trp Cys Glu Arg Thr Thr Leu
                               665
Ser Asn Gly Gly Cys Gln Tyr Leu Cys Leu Pro Ala Pro Gln Ile Asn
                           680
Pro His Ser Pro Lys Phe Thr Cys Ala Cys Pro Asp Gly Met Leu Leu
                                           700
                       695
Ala Arg Gly His Glu Glu Leu Pro His Arg Gly Leu Arg Leu Gln Trp
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                                       715
Pro Pro Arg Arg His Pro Pro Ser Gly
                725
                               729
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<210> 1256 <211> 264

<212> PRT

<213> Homo sapiens

<400> 1256

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<210> 1257 <211> 407 <212> PRT

<213> Homo sapiens

<400> 1257 Met Ser Gly Ala Pro Thr Ala Gly Ala Ala Leu Met Leu Cys Ala Ala 1 5 10 Thr Ala Val Leu Leu Ser Ala Gln Gly Pro Val Gln Ser Lys Ser 20 25 Pro Arg Phe Ala Ser Trp Asp Glu Met Asn Val Leu Ala His Gly Leu Leu Gln Leu Gly Gln Gly Leu Arg Glu His Ala Glu Arg Thr Arg Ser Gln Leu Ser Ala Leu Glu Arg Arg Leu Ser Ala Cys Gly Ser Ala Cys 75 Gln Gly Thr Glu Gly Ser Thr Asp Leu Pro Leu Ala Pro Glu Ser Arg 85 90 Val Asp Pro Glu Val Leu His Ser Leu Gln Thr Gln Leu Lys Ala Gln 105 Asn Ser Arg Ile Gln Gln Leu Phe His Lys Val Ala Gln Gln Gln Arg 120 His Leu Glu Lys Gln His Leu Arg Ile Gln His Leu Gln Ser Gln Phe 135 Gly Leu Leu Asp His Lys His Leu Asp His Glu Val Ala Lys Pro Ala 150 155 Arg Arg Lys Arg Leu Pro Glu Met Ala Gln Pro Val Asp Pro Ala His 165 170 Asn Val Ser Arg Leu His Arg Leu Pro Arg Asp Cys Gln Glu Leu Phe 185

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Gln Val Gly Glu Arg Gln Ser Gly Leu Phe Glu Ile Gln Pro Gln Gly
       195
                           200
Ser Pro Pro Phe Leu Val Asn Cys Lys Met Thr Ser Asp Gly Gly Trp
                        215
Thr Val Ile Gln Arg Arg His Asp Gly Ser Val Asp Phe Asn Arg Pro
                                       235
                    230
Trp Glu Ala Tyr Lys Ala Gly Phe Gly Asp Pro His Gly Glu Phe Trp
                245
                                   250
Leu Gly Leu Glu Lys Val His Ser Ile Thr Gly Asp Arg Asn Ser Arg
                               265
Leu Ala Val Gln Leu Arg Asp Trp Asp Gly Asn Ala Glu Leu Leu Gln
                           280
                                               285
Phe Ser Val His Leu Gly Gly Glu Asp Thr Ala Tyr Ser Leu Gln Leu
                       295
                                          300
Thr Ala Pro Val Ala Gly Gln Leu Gly Ala Thr Thr Val Pro Pro Ser
                   310
                                      315
Gly Leu Ser Val Pro Phe Ser Thr Trp Asp Gln Asp His Asp Leu Arg
              325 330
Arg Asp Lys Asn Cys Ala Lys Ser Leu Ser Gly Gly Trp Trp Phe Gly
           340
                               345
Thr Cys Ser His Ser Asn Leu Asn Gly Gln Tyr Phe Arg Ser Ile Pro
                          360
                                              365
Gln Gln Arg Gln Lys Leu Lys Lys Gly Ile Phe Trp Lys Thr Trp Arg
                       375
Gly Arg Tyr Tyr Pro Leu Gln Ala Thr Thr Met Leu Ile Gln Pro Met
                   390
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Ala Ala Glu Ala Ala Ser *
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<210> 1258 <211> 120 <212> PRT

<213> Homo sapiens

<400> 1258 Met Met Thr Pro Lys Leu Met Ile Trp Leu Leu Gln Ala Lys Ser 10 Ser Ile Ser Met Leu Glu Lys Ser Ser Lys Cys Leu Gly Arg Cys Phe 2.0 25 Ser Ser Phe Ala Lys Asn Leu Val Met Ile Gln Ser Cys Val Ser Trp 40 Ala Leu Met Ser Glu Asn Phe Tyr Arg Thr Leu Met Leu Cys Thr Thr 55 Thr Leu Leu Pro Ser Thr Gln Glu Cys Val His Leu Pro Leu Gly Ala 70 75 Leu Met Gln Lys Arg Ala Lys Asp Ser Phe Cys Thr Thr Gln Arg 85 90 Glu Lys Asp Phe Arg Ile Leu Ser Leu Glu Ser Ser Lys Gln Trp His 100 105 Asn Lys Ser Met Ala Leu Lys * 115 119

<210> 1259 <211> 160

<212> PRT <213> Homo sapiens

<400> 1259 Met Val Cys Leu Arg Leu Pro Gly Gly Ser Cys Met Ala Val Leu Thr 10 Val Thr Leu Met Val Leu Ser Ser Pro Leu Ala Leu Ala Gly Asp Thr 25 Arg Pro Arg Phe Leu Glu Tyr Ser Thr Gly Glu Cys Tyr Phe Phe Asn 40 Gly Thr Glu Arg Val Arg Phe Leu Asp Arg Tyr Phe Tyr Asn Gln Glu 55 60 Glu Tyr Val Arg Phe Asp Ser Asp Val Gly Glu Tyr Arg Ala Val Thr 70 Glu Leu Gly Arg Pro Asp Ala Glu Tyr Leu Glu Gln Pro Glu Gly Arg 90 Pro Trp Asn Ser Gln Lys Asp Ile Leu Glu Asp Glu Arg Ala Ala Val 105 Asp Thr Tyr Cys Arg His Asn Tyr Gly Val Val Glu Ser Phe Thr Val 120 Gln Arg Arg Val His Pro Lys Val Thr Val Tyr Pro Ser Lys Thr Gln 135 140 Pro Leu Gln Ala Pro Gln Pro Ala Val Leu Phe Cys Glu Trp Phe * 155

<210> 1260 <211> 111 <212> PRT <213> Homo sapiens

<400> 1260

<210> 1261 <211> 123 <212> PRT <213> Homo sapiens

<400> 1261

Met Ile Pro Ala Arg Phe Ala Gly Val Leu Leu Ala Leu Ala Leu Ile 10 Leu Pro Gly Thr Leu Cys Ala Glu Gly Thr Arg Gly Arg Ser Ser Thr Ala Arg Cys Ser Leu Phe Gly Ser Asp Phe Val Asn Thr Phe Asp Gly 40 Ser Met Tyr Ser Phe Ala Gly Tyr Cys Ser Tyr Leu Leu Ala Gly Gly Cys Gln Lys Arg Ser Phe Ser Ile Ile Gly Asp Phe Gln Asn Gly Lys 70 75 Arg Val Ser Leu Ser Val Tyr Leu Gly Glu Phe Phe Asp Ile His Leu 85 90 Phe Val Asn Gly Thr Val Thr Gln Gly Asp Gln Arg Val Ser Met Pro 100 105 Tyr Ala Ser Lys Gly Leu Tyr Leu Glu Thr * 120 122

<210> 1262 <211> 737 <212> PRT <213> Homo sapiens

<400> 1262 Met Phe Pro Ala Gly Pro Pro Trp Pro Arg Val Arg Val Val Gln Val 1.0 Leu Trp Ala Leu Leu Ala Val Leu Leu Ala Ser Trp Arg Leu Trp Ala 25 Ile Lys Asp Phe Gln Glu Cys Thr Trp Gln Val Val Leu Asn Glu Phe 40 Lys Arg Val Gly Glu Ser Gly Val Ser Asp Ser Phe Phe Glu Glu Glu 55 60 Pro Val Asp Thr Val Ser Ser Leu Phe His Met Leu Val Asp Ser Pro 70 75 Ile Asp Pro Ser Glu Lys Tyr Leu Gly Phe Pro Tyr Tyr Leu Lys Ile 85 90 Asn Tyr Ser Cys Glu Glu Lys Pro Ser Glu Asp Leu Val Arg Met Gly 105 His Leu Thr Gly Leu Lys Pro Leu Val Leu Val Thr Phe Gln Ser Pro 120 Val Asn Phe Tyr Arg Trp Lys Ile Glu Gln Leu Gln Ile Gln Met Glu 135 Ala Pro Phe Arg Ser Lys Gly Gly Pro Gly Gly Gly Arg Asp 155 150 Arg Asn Leu Ala Gly Met Asn Ile Asn Gly Phe Leu Lys Arg Asp Arg 165 170 Asp Asn Asn Ile Gln Phe Thr Val Gly Glu Glu Leu Phe Asn Leu Met 185 Pro Gln Tyr Phe Val Gly Val Ser Ser Arg Pro Leu Trp His Thr Val 195 200 Asp Gln Ser Pro Val Leu Ile Leu Gly Gly Ile Pro Asn Glu Lys Tyr 215 220 Val Leu Met Thr Asp Thr Ser Phe Lys Asp Phe Ser Leu Val Glu Val 235 Asn Gly Val Gly Gln Met Leu Ser Ile Asp Ser Cys Trp Val Gly Ser 250

Phe Tyr Cys Pro His Ser Gly Phe Thr Ala Thr Ile Tyr Asp Thr Ile

			260					265					270		
Ala	Thr	Glu 275	Ser	Thr	Leu	Phe	Ile 280	Arg	Gln	Asn	Gln	Leu 285	Val	Tyr	Tyr
Phe	Thr 290	Gly	Thr	Tyr	Thr	Thr 295	Leu	Tyr	Glu	Arg	Asn 300	Arg	Gly	Ser	Gly
Glu 305	Cys	Ala	Val	Ala	Gly 310	Pro	Thr	Pro	Gly	Glu 315	Gly	Thr	Leu	Val	Asn 320
Pro	Ser	Thr	Glu	Gly 325	Ser	Trp	Ile	Arg	Val 330	Leu	Ala	Ser	Glu	Cys 335	Ile
			Cys 340					345					350		
		355	Thr				360					365			
	370		Thr			375					380				_
385			Leu		390					395					400
			Gln	405					410				_	415	
			His 420					425					430		
		435	Ile				440					445			
	450		Ile Leu			455		_			460				_
465	ASII	FIIC	Бец	nea	470	ser	261	ASII	пуѕ	475	ASII	Pile	тте	1 7 1	480
			Pro	485					490	_			_	495	
			Val 500					505					510		
		515	Ser				520					525			
	530		Ile			535					540			_	
545	ASII	GIU	Thr	Met	ьеи 550	Thr	ьeu	Pne	Tyr	555	Asp	ser	гÀг	ьeu	1yr 560
			Tyr	565					570					575	
			Val 580					585					590		
		595	Glu				600					605			
	610		Phe			615					620				
625	Tyr	Thr	Arg	Gin	630	Arg	Tyr	Arg	Ala	Arg 635	Pro	Pro	Arg	Val	ьеи 640
Glu	Arg	Ser	Gly	Phe 645	Pro	Gln	Gly	Glu	Leu 650	Ala	Arg	His	Leu	Pro 655	Gly
Pro	Gly	Leu	Leu 660	Pro	Ala	Val	Ala	Ala 665	Leu	Arg	Val	Arg	Gln 670	Ala	Val
		675	Gly				680					685			
	690		Gly			695			_		700			_	_
705			Val		710					715					720
Arg	Val	Arg	Val	Arg 725	Ala	Ala	Gly	Ala	His 730	Phe	Pro	Gly	Gln	Gly 735	

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<210> 1266 <211> 148

<212> PRT <213> Homo sapiens

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<210> 1267 <211> 227 <212> PRT <213> Homo sapiens

<400> 1267 Arg Trp Leu Trp

Met Arg Trp Leu Trp Pro Leu Ala Val Ser Leu Ala Val Ile Leu Ala 1 5 10 Val Gly Leu Ser Arg Val Ser Gly Gly Ala Pro Leu His Leu Gly Arg 25 His Arg Ala Glu Thr Gln Glu Gln Gln Ser Arg Ser Lys Arg Gly Thr Glu Asp Glu Glu Ala Lys Gly Val Gln Gln Tyr Val Pro Glu Glu Trp 55 Ala Glu Tyr Pro Arg Pro Ile His Pro Ala Gly Leu Gln Pro Thr Lys 70 75 Pro Leu Val Ala Thr Ser Pro Asn Pro Asp Lys Asp Gly Gly Thr Pro 90 Asp Ser Gly Gln Glu Leu Arg Gly Asn Leu Thr Gly Ala Pro Gly Gln 105 Arg Leu Gln Ile Gln Asn Pro Leu Tyr Pro Val Thr Glu Ser Ser Tyr 120 125 Ser Ala Tyr Ala Ile Met Leu Leu Ala Leu Val Glu Phe Ala Ala Gly 135 140 Ile Val Gly Asn Leu Ser Val Met Cys Ile Ala Trp His Ser Tyr Tyr 150 155 Leu Lys Ser Ala Trp Asn Ser Ile Leu Ala Ser Leu Ala Leu Trp Asp 165 170 Phe Leu Val Leu Phe Phe Cys Leu Pro Ile Val Ile Leu Asn Glu Ile

Thr Lys Gln Arg Leu Leu Gly Asp Ala Pro Cys Pro Cys Arg Ala Leu
195 200 205

His Gly Gly Leu Leu Ser Gly Ser His Asp Phe Gln Pro Leu Cys Pro
210 215 220

Gly His *
225 226

<210> 1268 <211> 983 <212> PRT <213> Homo sapiens

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Pro Pro Gly Gly Ala Glu Ser Val His Ser Phe Tyr His Ala Asp Cys

			340					345					350		
His	Leu	Glu 355	Pro	Val	Arg	Cys	Gln 360	Ala	Pro	Pro	Pro	Arg 365	Ser	Pro	Ser
Glu	Ala 370	Ser	Gly	Arg	Thr	Val 375	Gly	Ser	Glγ	Lys	Val 380	Tyr	Pro	Thr	Val
His 385	Thr	Ser	Pro	Pro	Pro 390	Glu	Thr	Leu	Lys	Glu 395	Lys	Ala	Leu	Val	Glu 400
Val	Ala	Ala	Ser	Ser 405	Gly	Pro	Pro	Thr	Leu 410	Thr	Ser	Leu	Asn	Ile 415	Pro
Pro	Gly	Pro	Tyr 420	Ser	Ser	Met	His	Lys 425	Leu	Leu	Glu	Thr	Gln 430	Ser	Thr
Gly	Ala	Cys 435	Gln	Ser	Ser	Cys	Lys 440	Ile	Ser	Ser	Pro	Cys 445	Leu	Lys	Ala
Asp	Ser 450	Gly	Ala	Cys	Gly	Pro 455	Asp	Ser	Cys	Pro	Tyr 460	Cys	Ala	Arg	Ala
Gly 465	Ala	Gly	Glu	Val	Glu 470	Leu	Ala	Asp	Arg	Glu 475	Met	Pro	Asp	Ser	Asp 480
Ser	Glu	Ala	Val	Tyr 485	Glu	Phe	Thr	Gln	Asp 490	Ala	Gln	His	Ser	Asp 495	Leu
Arg	Asp	Pro	His 500	Ser	Arg	Arg	Gln	Arg 505	Ser	Leu	Gly	Pro	Asp 510	Ala	Glu
Pro	Ser	Ser 515	Val	Leu	Ala	Phe	Trp 520	Arg	Leu	Ile	Cys	Asp 525	Thr	Phe	Arg
Lys	Ile 530	Val	Asp	Ser	Lys	Tyr 535	Phe	Gly	Arg	Gly	Ile 540	Met	Ile	Ala	Ile
Leu 545	Val	Asn	Thr	Leu	Ser 550	Met	Gly	Ile	Glu	Tyr 555	His	Glu	Gln	Pro	Glu 560
Glu	Leu	Thr	Asn	Ala 565	Leu	Glu	Ile	Ser	Asn 570	Ile	Val	Phe	Thr	Ser 575	Leu
			Glu 580					585			_		590		
		59 5	Asn				600					605			
Ser	Val 610	Trp	Glu	Ile	Val	Gly 615	Gln	Gln	Gly	Gly	Gly 620	Leu	Ser	Val	Leu
Arg 625	Thr	Phe	Arg	Leu	Met 630	Arg	Val	Leu	Lys	Leu 635	Val	Arg	Phe	Leu	Pro 640
			Arg	645					650	-				655	
			Cys 660					665					670		
Leu	Gly	Met 675	His	Leu	Phe	Gly	Cys 680	Lys	Phe	Ala	Ser	Glu 685	Arg	Asp	Gly
	690		Pro			695					700				
Val 705	Thr	Val	Phe	Gln	Ile 710	Leu	Thr	Gln	Glu	Asp 715	Trp	Asn	Lys	Val	Leu 720
Tyr	Asn	Gly	Met	Ala 725	Ser	Thr	Ser	Ser	Trp 730	Ala	Ala	Leu	Tyr	Phe 735	Ile
			Thr 740				-	745					750		
		755	Glu				760					765			
	770		Asp			775				-	780	-	_		_
785			Leu		790				_	795					800
Lys	Ser	Leu	Leu	Pro 805	Pro	Leu	Ile	Ile	His 810	Thr	Ala	Ala	Thr	Pro 815	Met

Ser Leu Pro Lys Ser Thr Ser Thr Gly Leu Gly Glu Ala Leu Gly Pro 820 825 Ala Ser Arg Arg Thr Ser Ser Ser Gly Ser Ala Glu Pro Gly Ala Ala 840 845 835 His Glu Met Lys Ser Pro Pro Ser Ala Arg Ser Ser Pro His Ser Pro 860 855 Trp Ser Ala Ala Ser Ser Trp Thr Ser Arg Arg Ser Ser Arg Asn Ser 870 875 Leu Gly Arg Ala Pro Ser Leu Lys Arg Arg Ser Pro Ser Gly Glu Arg 890 Arg Ser Leu Leu Ser Gly Glu Gly Gln Glu Ser Gln Asp Glu Glu Glu 905 Ser Ser Glu Glu Glu Arg Ala Ser Pro Ala Gly Ser Asp His Arg His 920 Arg Gly Ser Leu Glu Arg Glu Ala Lys Ser Ser Phe Asp Leu Pro Asp 935 Thr Leu Gln Val Pro Gly Leu His Arg Thr Ala Ser Gly Arg Gly Ser 955 950 Ala Ser Glu His Gln Gly Leu Gln Trp Gln Val Gly Phe Arg Ala Pro 970 965 Gly Pro Gly Pro Ala Ala * 980 982

<210> 1269 <211> 708 <212> PRT <213> Homo sapiens

<400> 1269 Met Leu Ser Leu Arg Arg Cys Thr Ser Met Arg Leu Cys Leu Ser Ser 10 Ser Leu Ala Ser Pro Cys Ser Thr Met Leu Ser Thr Val Val Leu Tyr 20 25 Lys Val Cys Asn Ser Phe Val Glu Met Gly Ser Ala Asn Val Gln Ala 40 Thr Asp Tyr Leu Lys Gly Val Ala Ser Leu Phe Val Val Ser Leu Gly 55 Gly Ala Ala Val Gly Leu Val Phe Ala Phe Leu Leu Ala Leu Thr Thr 75 Arg Phe Thr Lys Arg Val Arg Ile Ile Glu Pro Leu Leu Val Phe Leu 90 85 Leu Ala Tyr Ala Ala Tyr Leu Thr Ala Glu Met Ala Ser Leu Ser Ala 105 Ile Leu Ala Val Thr Met Cys Gly Leu Gly Cys Lys Lys Tyr Val Glu 125 120 Ala Asn Ile Ser His Lys Ser Arg Thr Thr Val Lys Tyr Thr Met Lys 140 135 Thr Leu Ala Ser Cys Ala Glu Thr Val Ile Phe Met Leu Leu Gly Ile 155 150 Ser Thr Val Asp Ser Ser Lys Trp Ala Trp Asp Ser Gly Leu Val Leu 170 Gly Thr Leu Ile Phe Ile Leu Phe Phe Arg Ala Leu Gly Val Val Leu 185 Gln Thr Trp Val Leu Asn Gln Phe Arg Leu Val Pro Leu Asp Lys Ile 200 Asp Gln Val Val Met Ser Tyr Gly Gly Leu Arg Gly Ala Val Ala Phe

	210					215					220				
Ala 225	Leu	Val	Ile	Leu	Leu 230	Asp	Arg	Thr	Lys	Val 235	Pro	Ala	Lys	Asp	Tyr 240
	Val	Ala	Thr	Thr 245	Ile	Val	Val	Val	Phe 250	Phe	Thr	Val	Ile	Val 255	Gln
Gly	Leu	Thr	Ile 260	Lys	Pro	Leu	Val	Lys 265	Trp	Leu	Lys	Val	Lys 270	Arg	Ser
Glu	His	His 275	Lys	Pro	Thr	Leu	Asn 280	Gln	Glu	Leu	His	Glu 285	His	Thr	Phe
Asp	His 290	Ile	Leu	Ala	Ala	Val 295	Glu	Asp	Val	Val	Gly 300	His	His	Gly	Tyr
His 305	Tyr	Trp	Arg	Asp	Arg 310	Trp	Glu	Gln	Phe	Asp 315	Lys	Lys	Tyr	Leu	Ser 320
Gln	Leu	Leu	Met	Arg 325	Arg	Ser	Ala	Tyr	Arg 330	Ile	Arg	Asp	Gln	Ile 335	Trp
Asp	Val	Tyr	Tyr 340	Arg	Leu	Asn	Ile	Arg 345	Asp	Ala	Ile	Ser	Phe 350	Val	Asp
Gln	Gly	Gly 355	His	Val	Leu	Ser	Ser 360	Thr	Gly	Leu	Thr	Leu 365	Pro	Ser	Met
	370	_				375		Thr			380				
385					390			Asp		395					400
		-		405	_		_	Ala	410					415	
	_		420					Arg 425	_				430		
		435			_		440	Glu				445			
	450			_	_	455		Glu -			460				
465		_			470		-	Pro		475					480
_	_	=	_	485				Ala	490					495	
			500					Ala 505					510		
		515					520	Asp				525			
	530		_			535		Ala	_		540				
545					550			Leu		555					560
				565				Ala	570					575	
			580					Tyr 585					590		
		595					600	Gly				605			
	610					615		Cys			620				
625					630			Thr		635		•			640
				645		_		Ser	650		•			655	
			660	_		_		Glu 665					670		
Gln	Gln	Glu 675	Leu	Gln	Pro	Leu	Met 680	Gly	His	Lys	Asp	His 685	Thr	His	Leu

Ser Pro Gly Thr Ala Thr Ser His Trp Cys Ile Gln Phe Asn Arg Gly
690 695 700

Ser Arg Leu *
705 707

<210> 1270 <211> 93 <212> PRT <213> Homo sapiens

<210> 1271 <211> 648 <212> PRT <213> Homo sapiens

<400> 1271 Met Leu Trp Val Thr Gly Pro Val Leu Ala Val Ile Leu Ile Ile Leu 10 Ile Val Ile Ala Ile Leu Leu Phe Lys Arg Lys Arg Thr His Ser Pro 25 20 Ser Ser Lys Asp Glu Gln Ser Ile Gly Leu Lys Asp Ser Leu Leu Ala 35 40 His Ser Ser Asp Pro Val Glu Met Arg Arg Leu Asn Tyr Gln Thr Pro 55 Gly Met Arg Asp His Pro Pro Ile Pro Ile Thr Asp Leu Ala Asp Asn 70 75 Ile Glu Arg Leu Lys Ala Asn Asp Gly Leu Lys Phe Ser Gln Glu Tyr 85 Glu Ser Ile Asp Pro Gly Gln Gln Phe Thr Trp Glu Asn Ser Asn Leu 105 Glu Val Asn Lys Pro Lys Asn Arg Tyr Ala Asn Val Ile Ala Tyr Asp 115 120 His Ser Arg Val Ile Leu Thr Ser Ile Asp Gly Val Pro Gly Ser Asp 140 135 Tyr Ile Asn Ala Asn Tyr Ile Asp Gly Tyr Arg Lys Gln Asn Ala Tyr 155 150 Ile Ala Thr Gln Gly Pro Leu Pro Glu Thr Met Gly Asp Phe Trp Arg 170 165 Met Val Trp Glu Gln Arg Thr Ala Thr Val Val Met Met Thr Arg Leu

			180					185					190		
Glu	Glu	Lys 195	Ser	Arg	Val	Lys	Cys 200	Asp	Gln	Tyr	Trp	Pro 205	Ala	Arg	Gly
Thr	Glu 210	Thr	Cys	Gly	Leu	Ile 215	Gln	Val	Thr	Leu	Leu 220	Asp	Thr	Val	Glu
Leu 225	Ala	Thr	Tyr	Thr	Val 230	Arg	Thr	Phe	Ala	Leu 235	His	Lys	Ser	Gly	Ser 240
Ser	Glu	Lys	Arg	Glu 245	Leu	Arg	Gln	Phe	Gln 250	Phe	Met	Ala	Trp	Pro 255	Asp
His	Gly	Val	Pro 260	Glu	Tyr	Pro	Thr	Pro 265	Ile	Leu	Ala	Phe	Leu 270	Arg	Arg
Val	Lys	Ala 275	Cys	Asn	Pro	Leu	Asp 280	Ala	Gly	Pro	Met	Val 285	Val	His	Суѕ
Ser	Ala 290	Gly	Val	Gly	Arg	Thr 295	Gly	Cys	Phe	Ile	Val 300	Ile	Asp	Ala	Met
305			Met		310		_			315		_	_		320
			Arg	325					330					335	
Tyr	Val	Phe	Ile 340	His	Glu	Ala	Leu	Leu 345	Glu	Ala	Ala	Thr	Cys 350	Gly	His
Thr	Glu	Val 355	Pro	Ala	Arg	Asn	Leu 360	Tyr	Ala	His	Ile	Gln 365	Lys	Leu	Gly
	370		Pro			375					380				-
385			Ser		390		•			395					400
			Asn	405					410					415	_
			Arg 420					425		_			430		
		435	Asn				440					445			
	450		Thr			455					460				
465			Trp		470					475					480
			Met	485			_	_	490		_	_		495	
			Arg 500					505					510		_
		515	Gln				520					525			
	530		Ser			535	_				540		_	_	
545			Val		550					555					560
			Lys	565					570		_	_		575	
			Ser 580					585					590		
		595	Leu				600					605			
	610		Lys			615					620				
625			Tyr		630			Arg	Ala	Ala 635	Leu	Glu	Tyr	Leu	Gly 640
Ser	Phe	Asp	His	Tyr 645	Ala	Thr 647	*								

<210> 1272 <211> 109 <212> PRT <213> Homo sapiens

<400> 1272 Met Lys Ala Leu Cys Leu Leu Leu Pro Val Leu Gly Leu Leu Val 10 Ser Ser Lys Thr Leu Cys Ser Met Glu Glu Ala Ile Asn Glu Arg Ile 25 2.0 Gln Glu Val Ala Gly Ser Leu Ile Phe Arg Ala Ile Ser Ser Ile Gly 40 35 Leu Glu Cys Gln Ser Val Thr Ser Arg Gly Asp Leu Ala Thr Cys Pro 55 Arg Gly Phe Ala Val Thr Gly Cys Thr Cys Gly Ser Ala Cys Gly Ser 70 Trp Asp Val Arg Ala Glu Thr Thr Cys His Cys Gln Cys Ala Gly Met 90 85 Asp Trp Thr Gly Ala Arg Cys Cys Arg Val Gln Pro * 105 100

<210> 1273 <211> 56 <212> PRT <213> Homo sapiens

<210> 1274 <211> 188 <212> PRT <213> Homo sapiens

 400> 1274

 Met Asp Leu Ser Leu Leu Trp Val Leu Leu Pro Leu Val Thr Met Ala

 1
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 Trp Gly Gln Tyr Gly Asp Tyr Gly Tyr Pro Tyr Gln Gln Tyr His Asp

 20
 25
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 Tyr Ser Asp Asp Gly Trp Val Asn Leu Asn Arg Gln Gly Phe Ser Tyr

 35
 40
 45

 Gln Cys Pro Gln Gly Gln Val Ile Val Ala Val Arg Ser Ile Phe Ser

55 60 Lys Lys Glu Gly Ser Asp Arg Gln Trp Asn Tyr Ala Cys Met Pro Thr 70 75 Pro Gln Ser Leu Gly Glu Pro Thr Glu Cys Trp Trp Glu Glu Ile Asn 85 90 Arg Ala Gly Met Glu Trp Tyr Gln Thr Cys Ser Asn Asn Gly Leu Val 100 105 Ala Gly Phe Gln Ser Arg Tyr Phe Glu Ser Val Leu Asp Arg Glu Trp 115 120 125 Gln Phe Tyr Cys Cys Arg Tyr Ser Lys Arg Cys Pro Tyr Ser Cys Trp 135 Leu Thr Thr Glu Tyr Pro Gly His Tyr Gly Glu Glu Met Asp Met Ile 155 Ser Tyr Asn Tyr Asp Tyr Tyr Ile Arg Gly Ala Thr Thr His Phe Leu 165 170 Cys Ser Gly Lys Gly Ser Pro Ser Gly Ser Ser * 185

<210> 1275 <211> 81 <212> PRT

<213> Homo sapiens

<210> 1276 <211> 46 <212> PRT <213> Homo sapiens

<210> 1277

<211> 431 <212> PRT <213> Homo sapiens

<400> 1277 Met Ala Leu Leu Val Pro Leu Ala Leu Leu Val Ile Gln Ala His Leu 5 10 Val Leu Ser Val Gln Leu Glu Arg Val Val Thr Glu Glu Lys Val Ala 25 Leu Leu Ala Leu Leu Val Leu Pro Val Leu Leu Val Pro Glu Val Leu 40 Leu Val Leu Lys Ala His Val Val Thr Lys Val Lys Gln Val Asn Val Glu Leu Leu Ala Ser Lys Asp Ile Glu Asp Ser Leu Val Ile Gln Val 75 Pro Gln Val Leu Gln Ala Leu Leu Val Ser Arg Val Gln Ser Ala Val 85 90 Gln Asp Leu Gln Ala Pro Glu Asp Leu Leu Asp Pro Val Asp Leu Leu 100 105 Ala Lys Met Glu Pro Val Asp Ile Gln Val Pro Leu Asp His Gln Gly 125 120 Leu Glu Val Thr Glu Val Lys Glu Asp Leu Arg Ala Pro Gln Ala Thr 135 140 Gln Gly Asn Gln Ala Leu Leu Asp Leu Leu Val Pro Leu Val Leu Ala 150 155 Val Val Leu Glu Pro Leu Pro Leu Gly Leu Glu Val Lys Lys 165 170 Leu Ala Gly Phe Ala Pro Tyr Tyr Gly Asp Glu Pro Met Asp Phe Lys 180 185 190 Ile Asn Thr Asp Glu Ile Met Thr Ser Leu Lys Ser Val Asn Gly Gln 200 205 Ile Glu Ser Leu Ile Ser Pro Asp Gly Ser Arg Lys Asn Pro Ala Arg 215 220 Asn Cys Arg Asp Leu Lys Phe Cys His Pro Glu Leu Lys Ser Gly Glu 230 235 Tyr Trp Val Asp Pro Asn Gln Gly Cys Lys Leu Asp Ala Ile Lys Val 250 Phe Cys Asn Met Glu Thr Gly Glu Thr Cys Ile Ser Ala Asn Pro Leu 265 Asn Val Pro Arg Lys His Trp Trp Thr Asp Ser Ser Ala Glu Lys Lys 280 His Val Trp Phe Gly Glu Ser Met Asp Gly Gly Phe Gln Phe Ser Tyr 295 Gly Asn Pro Glu Leu Pro Glu Asp Val Leu Asp Val Gln Leu Ala Phe 315 310 Leu Arg Leu Leu Ser Ser Arg Ala Ser Gln Asn Ile Thr Tyr His Cys 330 325 Lys Asn Ser Ile Ala Tyr Met Asp Gln Ala Ser Gly Asn Val Lys Lys 345 Ala Leu Lys Leu Met Gly Ser Asn Glu Gly Glu Phe Lys Ala Glu Gly 360 Asn Ser Lys Phe Thr Tyr Thr Val Leu Glu Asp Gly Cys Thr Lys His 375 380 Thr Gly Glu Trp Ser Lys Thr Val Phe Glu Tyr Arg Thr Arg Lys Ala 390 395 Val Arg Leu Pro Ile Val Asp Ile Ala Pro Tyr Asp Ile Gly Gly Pro 410 Asp Gln Glu Phe Gly Val Asp Val Gly Pro Val Cys Phe Leu *

420 425 430

<210> 1278 <211> 53 <212> PRT

<213> Homo sapiens

<210> 1279 <211> 73 <212> PRT

<213> Homo sapiens

<210> 1280 <211> 51 <212> PRT <213> Homo sapiens

<210> 1281 <211> 144 <212> PRT <213> Homo sapiens

<400> 1281 Met Lys Ser Gly Ser Gly Gly Ser Pro Thr Ser Leu Trp Gly Leu 10 Leu Phe Leu Ser Ala Ala Leu Ser Leu Trp Pro Thr Ser Gly Glu Ile 20 25 Cys Gly Pro Gly Ile Asp Ile Arg Asn Asp Tyr Gln Gln Leu Lys Arg Leu Glu Asn Cys Thr Val Ile Glu Gly Tyr Leu His Ile Leu Leu Ile 55 Ser Lys Ala Glu Asp Tyr Arg Ser Tyr Arg Phe Pro Lys Leu Thr Val 70 75 Ile Thr Glu Tyr Leu Leu Phe Arg Val Ala Gly Leu Glu Ser Leu 85 90 Gly Asp Leu Phe Pro Asn Leu Thr Val Ile Arg Gly Trp Lys Leu Phe 105 Tyr Asn Tyr Ala Leu Val Ile Phe Glu Met Thr Asn Leu Lys Asp Ile 120 Gly Leu Tyr Asn Leu Arg Asn Ile Thr Arg Gly Gly His Gln Asp 135 140

<210> 1282 <211> 267 <212> PRT <213> Homo sapiens

<400> 1282 Met Gly Pro Pro Ser Ala Cys Pro His Arg Glu Cys Ile Pro Trp Gln 10 Gly Leu Leu Thr Ala Ser Leu Leu Thr Phe Trp Asn Ala Pro Thr 2.0 25 Thr Ala Trp Leu Phe Ile Ala Ser Ala Pro Phe Glu Val Ala Glu Gly Glu Asn Val His Leu Ser Val Val Tyr Leu Pro Glu Asn Leu Tyr Ser 55 Tyr Gly Trp Tyr Lys Gly Lys Thr Val Glu Pro Asn Gln Leu Ile Ala 75 Ala Tyr Val Ile Asp Asp Thr His Val Arg Thr Pro Gly Pro Ala Tyr 85 . 90 Ser Gly Arg Glu Thr Ile Ser Pro Ser Gly Asp Leu His Phe Gln Asn 105 110 Val Thr Leu Glu Asp Thr Gly Tyr Tyr Asn Leu Gln Val Thr Tyr Arg 120 125 Asn Ser Gln Ile Glu Gln Ala Ser His His Leu Arg Val Tyr Gln Val 135 140 Ser Gly Leu Thr Pro Pro Ser Lys Pro Ala Ala Pro Gln Ser Pro Arg 150 155 Arg Ala Pro Gly Val Leu Thr Cys His Thr Asn Asn Thr Gly Thr Ser 165 170 Phe Gln Trp Ile Phe Asn Asn Gln Arg Leu Gln Val Thr Lys Arg Met

<210> 1283 <211> 262 <212> PRT <213> Homo sapiens

<400> 1283

Met Leu Val Leu Val Leu Arg Val Ser Leu Ala Ala Leu Val Lys 1 5 10 Met Glu Leu Leu Val Arg Trp Ala Pro Val Ala Cys Leu Val Arg Glu 25 Val Ala Leu Glu Pro Leu Ala Leu Leu Val Leu Val Glu Met Met Val 40 Leu Leu Val Leu Pro Gly Pro Leu Val Pro Pro Ala Pro Leu Val Leu 55 Leu Ala Ser Leu Val Leu Leu Val Leu Arg Val Lys Leu Val Pro Lys 70 75 Gly Pro Glu Ala Leu Lys Val Pro Arg Val Cys Val Val Ser Leu Ala 90 Pro Leu Ala Leu Leu Val Leu Leu Ala Leu Leu Glu Thr Leu Val Leu 105 Arg Glu Ser Leu Val Leu Lys Val Pro Met Val Leu Leu Val Leu Leu 120 Val Leu Leu Ala Ser Leu Val Pro Glu Ala Pro Leu Asp Pro Arg Ala 135 Pro Ala Ala Leu Leu Val Pro Arg Val Thr Ala Val Asn Leu Val Leu 150 155 Leu Ala Ala Lys Glu Thr Leu Val Leu Arg Glu Ser Leu Ala Leu Leu 165 170 Val Phe Lys Asp Pro Leu Ala Leu Leu Glu Arg Lys Glu Ser Glu Glu 180 185 Leu Glu Val Asn Pro Asp Pro Leu Ala Cys Pro Asp Pro Leu Ala Ser 200 205 Val Val Asp Leu Val Ala Val Val Ser Leu Ala Gln Met Val Leu Leu 215 Val Pro Arg Val Pro Leu Val Asn Val Leu Leu Ala Leu Leu Ala 230 235 Pro Lys Asp Leu Leu Val Lys Leu Val Val Pro Val Lys Leu Val Cys 245 250 Leu Val Pro Arg Val * 260 261

<210> 1284

<211> 50 <212> PRT <213> Homo sapiens

<210> 1285 <211> 323 <212> PRT <213> Homo sapiens

<400> 1285 Met Leu Val Met Ala Pro Arg Thr Val Leu Leu Leu Ser Ala Ala 10 Leu Ala Leu Thr Glu Thr Trp Ala Gly Ser His Ser Met Arg Tyr Phe 25 Tyr Thr Ser Val Ser Arg Pro Gly Arg Gly Glu Pro Arg Phe Ile Ser 40 Val Gly Tyr Val Asp Asp Thr Gln Phe Val Arg Phe Asp Ser Asp Ala 55 Ala Ser Pro Arg Glu Glu Pro Arg Ala Pro Trp Ile Glu Gln Glu Gly 70 Pro Glu Tyr Trp Asp Arg Asn Thr Gln Ile Tyr Lys Ala Gln Ala Gln 85 90 Thr Asp Arg Glu Ser Leu Arg Asn Leu Arg Gly Tyr Tyr Asn Gln Ser 100 105 Glu Ala Gly Ser His Thr Leu Gln Ser Met Tyr Gly Cys Asp Val Gly 120 Pro Asp Gly Arg Leu Leu Arg Gly His Asp Gln Tyr Ala Tyr Asp Gly 135 140 Lys Asp Tyr Ile Ala Leu Asn Glu Asp Leu Arg Ser Trp Thr Ala Ala 150 155 Asp Thr Ala Ala Gln Ile Thr Gln Arg Lys Trp Glu Ala Ala Arg Glu 170 Ala Glu Gln Arg Arg Ala Tyr Leu Glu Gly Glu Cys Val Glu Trp Leu 180 Arg Arg Tyr Leu Glu Asn Gly Lys Asp Lys Leu Glu Arg Ala Asp Pro 200 Pro Lys Thr His Val Thr His His Pro Ile Ser Asp His Glu Ala Thr 215 220 Leu Arg Cys Trp Ala Leu Gly Phe Tyr Pro Ala Glu Ile Thr Leu Thr 230 235 Trp Gln Arg Asp Gly Glu Asp Gln Thr Gln Asp Thr Glu Leu Val Glu 250 Thr Arg Pro Ala Gly Asp Arg Thr Phe Gln Lys Val Gly Gln Leu Trp 265

Val Val Pro Ser Gly Glu Glu Gln Arg Tyr Thr Cys His Val Gln His

<210> 1286 <211> 306 <212> PRT <213> Homo sapiens

<400> 1286 Met Leu Leu Phe Leu Leu Ser Ala Leu Val Leu Leu Thr Gln Pro Leu 1 5 10 Gly Tyr Leu Glu Ala Glu Met Lys Thr Tyr Ser His Arg Thr Met Pro 2.5 Ser Ala Cys Thr Leu Val Met Cys Ser Ser Val Glu Ser Gly Leu Pro 40 Gly Arg Asp Gly Arg Asp Gly Arg Glu Gly Pro Arg Gly Glu Lys Gly 55 Asp Pro Gly Leu Pro Gly Ala Ala Gly Gln Ala Gly Met Pro Gly Gln 70 75 Ala Gly Pro Val Gly Pro Lys Gly Asp Asn Gly Ser Val Gly Glu Pro 90 Gly Pro Lys Gly Asp Thr Gly Pro Ser Gly Pro Pro Gly Pro Pro Gly 105 Val Pro Gly Pro Ala Gly Arg Glu Gly Pro Leu Gly Lys Gln Gly Asn 120 Ile Gly Pro Gln Gly Lys Pro Gly Pro Lys Gly Glu Ala Gly Pro Lys 135 Gly Glu Val Gly Ala Pro Gly Met Gln Gly Ser Ala Gly Ala Arg Gly 150 Leu Ala Gly Pro Lys Gly Glu Arg Gly Val Pro Gly Glu Arg Gly Val 165 170 Pro Gly Asn Thr Gly Ala Ala Gly Ser Ala Gly Ala Met Gly Pro Gln 185 Gly Ser Pro Gly Ala Arg Gly Pro Pro Gly Leu Lys Gly Asp Lys Gly 200 Ile Pro Gly Asp Lys Gly Ala Lys Gly Glu Ser Gly Leu Pro Asp Val 215 220 Ala Ser Leu Arg Gln Gln Val Glu Ala Leu Gln Gly Gln Val Gln His 230 235 Leu Glm Ala Ala Phe Ser Glm Tyr Lys Lys Val Glu Leu Phe Pro Asm 245 250 Gly Gln Ser Val Gly Glu Lys Ile Phe Lys Thr Ala Gly Phe Val Lys 260 265 Pro Phe Thr Glu Ala Gln Leu Leu Cys Thr Gln Ala Gly Gly Gln Leu 280 285 Ala Ser Pro Arg Ser Ala Ala Glu Asn Ala Pro Leu Ala Thr Ala Gly 295 300 Pro * 305

<210> 1287 <211> 299 <212> PRT <213> Homo sapiens

<400> 1287 Met Gly Arg Trp Ala Leu Asp Val Ala Phe Leu Trp Lys Ala Val Leu 5 10 Thr Leu Gly Leu Val Leu Leu Tyr Tyr Cys Phe Ser Ile Gly Ile Thr 20 25 Phe Tyr Asn Lys Trp Leu Thr Lys Ser Phe His Phe Pro Leu Phe Met 40 Thr Met Leu His Leu Ala Val Ile Phe Leu Phe Ser Ala Leu Ser Arg 55 Ala Leu Val Gln Cys Ser Ser His Arg Ala Arg Val Val Leu Ser Trp 75 Ala Asp Tyr Leu Arg Arg Val Ala Pro Thr Ala Leu Ala Thr Ala Leu 90 Asp Val Gly Leu Ser Asn Trp Ser Phe Leu Tyr Val Thr Val Ser Leu 105 Tyr Thr Met Thr Lys Ser Ser Ala Val Leu Phe Ile Leu Ile Phe Ser 120 125 Leu Ile Phe Lys Leu Glu Glu Leu Arg Ala Ala Leu Val Leu Val Val 135 140 Leu Leu Ile Ala Gly Gly Leu Phe Met Phe Thr Tyr Lys Ser Thr Gln 150 155 Phe Asn Val Glu Gly Phe Ala Leu Val Leu Gly Ala Ser Phe Ile Gly 165 170 Gly Ile Arg Trp Thr Leu Thr Gln Met Leu Leu Gln Lys Ala Glu Leu 185 Gly Leu Gln Asn Pro Ile Asp Thr Met Phe His Leu Gln Pro Leu Met 200 205 Phe Leu Gly Leu Phe Pro Leu Phe Ala Val Phe Glu Gly Leu His Leu 215 220 Ser Thr Ser Glu Lys Ile Phe Arg Phe Gln Gly His Arg Ala Ala Pro 230 235 Ala Gly Thr Trp Gly Ala Ser Ser Leu Ala Gly Phe Ser Pro Leu Val 245 250 Trp Ala Ser Leu Ser Ser Ser Trp Ser Pro Glu Pro Pro Ala Ser Leu 265 Ser Pro Leu Pro Ala Phe Leu Arg Lys Ser Ala Leu Cys Cys Trp Gln 280 Leu Ile Cys Trp Ala Ile Arg Ser Ala Ser 290 295

<210> 1288 <211> 161 <212> PRT <213> Homo sapiens

<400> 1288
Met Glu Ser Ala Leu Pro Ala Ala Gly Phe Leu Tyr Trp Val Gly Ala
1 5 10 15
Gly Thr Val Ala Tyr Leu Ala Leu Arg Ile Ser Tyr Ser Leu Phe Thr

25 Ala Leu Arg Val Trp Gly Val Gly Asn Glu Ala Gly Val Gly Pro Gly 35 40 Leu Gly Glu Trp Ala Val Val Thr Gly Ser Thr Asp Gly Ile Gly Lys 55 Ser Tyr Ala Glu Glu Leu Ala Lys His Gly Met Lys Val Val Leu Ile 70 75 Ser Arg Ser Lys Asp Lys Leu Asp Gln Val Ser Ser Glu Ile Lys Glu 85 90 Lys Phe Lys Val Glu Thr Arg Thr Ile Ala Val Asp Phe Ala Ser Glu 105 Asp Ile Tyr Asp Lys Ile Lys Thr Gly Leu Ala Gly Leu Glu Ile Gly 120 Ile Leu Val Asn Asn Val Gly Met Ser Tyr Glu Tyr Pro Glu Tyr Phe 135 140 Leu Asp Val Pro Asp Leu Asp Asn Val Ile Lys Lys Asn Asp Lys Tyr 150 155

<210> 1289 <211> 46 <212> PRT <213> Homo sapiens

<210> 1290 <211> 453 <212> PRT <213> Homo sapiens

 <400> 1290

 Met Thr Ser Lys
 Phe Ile Leu Val Ser Phe Ile Leu Ala Ala Leu Ser 10

 1
 5
 6

 Leu Ser Thr Thr Phe Ser Leu Gln Pro Asp Gln Gln Lys
 Val Leu Leu 20

 Val Ser Phe Asp Gly Phe Arg Trp Asp Tyr Leu Tyr Lys
 Val Pro Thr 35

 Pro His Phe His Tyr Ile Met Lys
 Tyr Gly Val His Val Lys
 Gln Val 55

 Thr Asn Val Phe Ile Thr Lys
 Thr Tyr Pro Asn His Tyr Thr Leu Val 65
 70
 75
 80

 Thr Gly Leu Phe Ala Glu Asn His Gly Ile Val Ala Asn Asp Met Phe 85
 90
 95

 Asp Pro Ile Arg Asn Lys Ser Phe Ser Leu Asp His Met Asn Ile Tyr 100
 Tyr 105
 100

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Asp Ser Lys Phe Trp Glu Glu Ala Thr Pro Ile Trp Ile Thr Asn Gln
    115
                        120
Arg Ala Gly His Thr Ser Gly Ala Ala Met Trp Pro Gly Thr Asp Val
                       135
Lys Ile His Lys Arg Phe Pro Thr His Tyr Met Pro Tyr Asn Glu Ser
                    150
                                       155
Val Ser Phe Glu Asp Arg Val Ala Lys Ile Ile Glu Trp Phe Thr Ser
                165
                                   170
Lys Glu Pro Ile Asn Leu Gly Leu Leu Tyr Trp Glu Asp Pro Asp Asp
                               185
Met Gly His His Leu Gly Pro Asp Ser Pro Leu Met Gly Pro Val Ile
                           200
                                               205
Ser Asp Ile Asp Lys Lys Leu Gly Tyr Leu Ile Gln Met Leu Lys Lys
                       215
                                           220
Ala Lys Leu Trp Asn Thr Leu Asn Leu Ile Ile Thr Ser Asp His Gly
                  230
                                       235
Met Thr Gln Cys Ser Glu Glu Arg Leu Ile Glu Leu Asp Gln Tyr Leu
               245
                                  250
Asp Lys Asp His Tyr Thr Leu Ile Asp Gln Ser Pro Val Ala Ala Ile
           260
                               265
Leu Pro Lys Glu Gly Lys Phe Asp Glu Val Tyr Glu Ala Leu Thr His
       275
                           280
Ala His Pro Asn Leu Thr Val Tyr Lys Lys Glu Asp Val Pro Glu Arg
                       295
                                           300
Trp His Tyr Lys Tyr Asn Ser Arg Ile Gln Pro Ile Ile Ala Val Ala
                   310
Asp Glu Gly Trp His Ile Leu Gln Asn Lys Ser Asp Asp Phe Leu Leu
               325
                                   330
Gly Asn His Gly Tyr His Asn Ala Leu Ala Asp Met His Pro Ile Phe
           340
                               345
Leu Ala His Gly Pro Ala Phe Arg Lys Asn Phe Ser Lys Glu Ala Met
                           360
Asn Ser Thr Asp Leu Tyr Pro Leu Leu Cys His Leu Leu Asn Ile Thr
                       375
                                           380
Ala Met Pro His Asn Gly Ser Phe Trp Asn Val Gln Asp Leu Leu Asn
                   390
                                       395
Ser Ala Met Pro Arg Val Val Pro Tyr Thr Gln Ser Thr Ile Leu Leu
               405
                                   410
Pro Gly Ser Val Lys Pro Ala Glu Tyr Asp Gln Glu Gly Ser Tyr Pro
                               425
                                                   430
Tyr Phe Ile Gly Val Ser Leu Gly Ser Ile Ile Val Ile Val Phe Phe
       435
                           440
Cys Asn Phe His *
    450
          452
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<210> 1291

<211> 78

<212> PRT

<213> Homo sapiens

<221> misc_feature

<222> (1)...(78)

<223> Xaa = any amino acid or nothing
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<400> 1291
Met Leu Ser Val Thr Ala Phe Ile Leu Ala Glu Thr Val Leu Ala Ser

<210> 1292 <211> 416 <212> PRT <213> Homo sapiens

<400> 1292 Met Val Leu Trp Ile Leu Trp Arg Pro Phe Gly Phe Ser Gly Arg Phe 1 5 10 Leu Lys Leu Glu Ser His Ser Ile Thr Glu Ser Lys Ser Leu Ile Pro 20 Val Ala Trp Thr Ser Leu Thr Gln Met Leu Leu Glu Ala Pro Gly Ile Phe Leu Leu Gly Gln Arg Lys Arg Phe Ser Thr Met Pro Glu Thr Glu 60 Thr His Glu Arg Glu Thr Glu Leu Phe Ser Pro Pro Ser Asp Val Arg 75 Gly Met Thr Lys Leu Asp Arg Thr Ala Phe Lys Lys Thr Val Asn Ile 90 Pro Val Leu Lys Val Arg Lys Glu Ile Val Ser Lys Leu Met Arg Ser 105 Leu Lys Arg Ala Ala Leu Gln Arg Pro Gly Ile Arg Arg Val Ile Glu 120 125 Asp Pro Glu Asp Lys Glu Ser Arg Leu Ile Met Leu Asp Pro Tyr Lys 135 Ile Phe Thr His Asp Ser Phe Glu Lys Ala Glu Leu Ser Val Leu Glu 150 155 160 Gln Leu Asn Val Ser Pro Gln Ile Ser Lys Tyr Asn Leu Glu Leu Thr 165 170 Tyr Glu His Phe Lys Ser Glu Glu Ile Leu Arg Ala Val Leu Pro Glu 185 Gly Gln Asp Val Thr Ser Gly Phe Ser Arg Ile Gly His Ile Ala His 200 205 Leu Asn Leu Arg Asp His Gln Leu Pro Phe Lys His Leu Ile Gly Gln 215 220 Val Met Ile Asp Lys Asn Pro Gly Ile Thr Ser Ala Val Asn Lys Ile 230 235 Asn Asn Ile Asp Asn Met Tyr Arg Asn Phe Gln Met Glu Val Leu Ser 250 Gly Glu Gln Asn Met Met Thr Lys Val Arg Glu Asn Asn Tyr Thr Tyr 265 Glu Phe Asp Phe Ser Lys Val Tyr Trp Asn Pro Arg Leu Ser Thr Glu 280 His Ser Arg Ile Thr Glu Leu Lys Pro Gly Asp Val Leu Phe Asp 295 300 Val Phe Ala Gly Val Gly Pro Phe Ala Ile Pro Val Ala Lys Lys Asn 310 315

Cys Thr Val Phe Ala Asn Asp Leu Asn Pro Glu Ser His Lys Trp Leu 330 325 Leu Tyr Asn Cys Lys Leu Asn Lys Val Asp Gln Lys Val Lys Val Phe 340 345 Asn Leu Asp Gly Lys Asp Phe Leu Gln Gly Pro Val Lys Glu Glu Leu 360 365 Met Gln Leu Leu Gly Leu Ser Lys Glu Arg Lys Pro Ser Val His Val 375 380 Val Met Asn Leu Pro Ala Lys Ala Ile Glu Phe Leu Ser Ala Phe Lys 390 395 Trp Leu Leu Asp Gly Gln Pro Met Pro Ala Val Ser Ser Phe Pro * 405 410

<210> 1293

<211> 113

<212> PRT

<213> Homo sapiens

<400> 1293

Met Val Arg Pro Leu Leu Leu Leu Asn Leu His Phe His Leu Pro Ser 10 Leu Val Ser Leu Ser Leu Leu Leu Ser Val Ser Leu Ser Leu 2.0 25 Val Asn Ala Val Arg Leu Leu Arg Ala Ser Phe Cys Ser Trp Leu Ile 40 Ala Lys Ser Leu Ile Thr Leu Trp Val Arg Pro Ser Gln Ile Gly Lys 55 Leu Lys Ala Leu Ala Ser Ser Thr Thr Ser Met Ala Trp Glu Gly Leu 70 75 Leu Asp Thr Phe Ala Leu Ser Ile Ser Ser Phe Ser Asn Ser Leu Leu 85 90 Gly Ile Leu Leu Cys Phe Leu Lys Ser Pro Asn Ile Phe Gln Ala Ser 100 105 110 112

<210> 1294

<211> 57

<212> PRT

<213> Homo sapiens

<400> 1294

 Met
 Asp
 Phe
 Leu
 Met
 Leu
 Ala
 Val
 Cys
 Ala
 His
 Arg
 Leu
 Cys
 Phe
 Leu

 Tyr
 Leu
 Phe
 Ile
 Leu
 Tyr
 Glu
 Ser
 Lys
 Asn
 Lys
 Arg
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<210> 1295 <211> 68 <212> PRT <213> Homo sapiens

<210> 1296 <211> 66 <212> PRT <213> Homo sapiens

<210> 1297 <211> 57 <212> PRT <213> Homo sapiens

<210> 1298

<211> 235 <212> PRT <213> Homo sapiens

<400> 1298 Met Arg Lys Thr Arg Leu Trp Gly Leu Leu Trp Met Leu Phe Val Ser 10 Glu Leu Arg Ala Ala Thr Lys Leu Thr Glu Glu Lys Tyr Glu Leu Lys 20 25 Glu Gly Gln Thr Leu Asp Val Lys Cys Asp Tyr Thr Leu Glu Lys Phe 40 Ala Ser Ser Gln Lys Ala Trp Gln Ile Ile Arg Asp Gly Glu Met Pro 55 Lys Thr Leu Ala Cys Thr Glu Arg Pro Ser Lys Asn Ser His Pro Val 70 75 Gln Val Gly Arg Ile Ile Leu Glu Asp Tyr His Asp His Gly Leu Leu 90 Arg Val Arg Met Val Asn Leu Gln Val Glu Asp Ser Gly Leu Tyr Gln 105 Cys Val Ile Tyr Gln Pro Pro Lys Glu Pro His Met Leu Phe Asp Arg 120 Ile Arg Leu Val Val Thr Lys Gly Phe Ser Gly Thr Pro Gly Ser Asn 135 140 Glu Asn Ser Thr Gln Asn Val Tyr Lys Ile Pro Pro Thr Thr Lys 150 155 Ala Leu Cys Pro Leu Tyr Thr Thr Pro Arg Thr Val Thr Gln Ala Pro 165 170 Pro Lys Ser Thr Ala Asp Val Ser Thr Pro Asp Ser Glu Ile Asn Leu 185 Thr Asn Val Thr Asp Ile Ile Arg Val Pro Val Phe Asn Ile Val Ile 200 Leu Leu Ala Gly Gly Phe Leu Ser Lys Ser Leu Val Phe Ser Val Leu 215 Phe Ala Val Thr Leu Arg Ser Phe Val Pro

<210> 1299 <211> 64 <212> PRT <213> Homo sapiens

230

<210> 1300 <211> 80

<212> PRT <213> Homo sapiens

<210> 1301 <211> 87 <212> PRT <213> Homo sapiens

<210> 1302 <211> 143 <212> PRT <213> Homo sapiens

 Val
 Pro
 Gln
 Ala
 Gly
 Gln
 His
 Ala
 Arg
 Gly
 Gln
 His
 Ala
 Met
 Gln

 Phe
 Pro
 Ala
 Glu
 Leu
 Thr
 Arg
 Asp
 Ala
 Cys
 Lys
 Thr
 Arg
 Pro
 Arg
 Glu

 Leu
 Arg
 Leu
 Ile
 Cys
 Ile
 Tyr
 Phe
 Ser
 Asn
 Thr
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 Phe
 Phe
 Lys

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<210> 1303 <211> 60 <212> PRT <213> Homo sapiens

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<210> 1306 <211> 138 <212> PRT

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<210> 1307 <211> 64 <212> PRT <213> Homo sapiens

<210> 1308 <211> 65 <212> PRT <213> Homo sapiens

<400> 1308

 Met
 Pro
 Cys
 Ser
 Gly
 Ser
 Val
 Gln
 Thr
 Phe
 Arg
 Pro
 Leu
 Leu
 Ile

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 Asn
 Val
 Thr
 Phe
 Phe
 Phe
 Ile
 Leu
 Pro
 Val
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 Cys
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 Asn
 Ala

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 Ile
 Asn
 Val
 Leu
 Glu
 Arg
 Pro
 Phe
 Trp
 Gln
 Leu
 Leu
 Gly
 Glu
 Ile

 Gly
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 Trp
 Leu
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<210> 1309 <211> 75 <212> PRT <213> Homo sapiens

<210> 1310 <211> 46 <212> PRT <213> Homo sapiens

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<210> 1311 <211> 105 <212> PRT <213> Homo sapiens

<210> 1312 <211> 114 <212> PRT <213> Homo sapiens

<400> 1312

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 Gly
 Lys
 Trp
 Cys
 Cys
 Ser
 Leu
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 Cys
 Gln
 Ser
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 Ser
 Leu
 Gly
 Pro
 Pro
 Gly

 Gln
 Thr
 Ala
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 Val
 Cys
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<210> 1314 <211> 65 <212> PRT <213> Homo sapiens

<400> 1314

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 Thr
 Val
 Val
 Ser
 Ser
 Arg
 Pro
 Val
 Ala
 Cys
 Leu
 Glu
 Ser
 Val

 Pro
 Gly
 Met
 Cys
 Met
 Cys
 Met
 Pro
 Leu
 Asn
 Tyr
 Arg
 Gly
 Ser

 Asn
 Phe
 Ser
 Glu
 Thr
 Asp
 Val
 Trp
 Met
 Asp
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 Leu

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<210> 1315 <211> 71 <212> PRT <213> Homo sapiens

<210> 1316 <211> 114 <212> PRT <213> Homo sapiens

65 70 75 80

Gly Leu Ala Ala Leu Pro Gly Ser Gly Ala Phe Ser Val Ile Pro Val
85 90 95

Ser Leu Leu Leu Pro Val Pro Glu Gly Leu Gly Arg Thr Tyr Leu Tyr
100 105 110

Ser *

113

<210> 1317 <211> 91 <212> PRT <213> Homo sapiens

(213) Homo Sapiens

<210> 1318 <211> 65 <212> PRT <213> Homo sapiens

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<211> 55

<212> PRT

<213> Homo sapiens

<400> 1321

<210> 1322

<211> 301

<212> PRT

<213> Homo sapiens

<400> 1322

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 Lys
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 Gly
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 Leu
 Trp
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 Glu
 Ile
 Leu
 Lys
 Leu
 Lys
 Leu
 Lys
 Lys
 Lys
 His
 Trp

 Pro
 Pro
 Trp
 Thr
 Leu
 Leu
 His
 Leu
 Glu
 Cys
 Phe
 Lys
 Lys
 His
 Trp

 Leu
 Ala
 Val
 Phe
 Glu
 Leu
 Val
 Met
 Glu
 Lys
 Asn
 Leu
 Leu
 Thr
 Ile
 Ile
 Leu
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 Phe

 Glu
 Ser
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 Tyr
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 Asn
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 Ala
 Asn
 Lys
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 Val
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 Ala
 Ile
 Ala
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 Lys
 Ala
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70
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Phe Ser Thr Arg Ser Asn Tyr Asp Gly Ile Leu Pro Gln Thr Phe Ala
             85
                                90
Gln Val Asn Asn Leu Leu Gln Thr Phe Ala Glu Val Lys Thr Lys Leu
     100 105 110
Lys Pro Asn Ser Ser Glu Asn Thr Val Thr Lys Lys Gln Glu Gly Thr
 115 120
Ser Leu Lys Asn Ser His Asn Gln Glu Ile Thr Val Phe Ser Ser Ser
                    135
His Leu Pro Gln Pro Ser Arg His Gln Glu Ile Trp Ser Ile Leu Glu
                                   155
Ser Val Trp Ile Thr Ile Tyr Gln Asn Ser Thr Asp Val Phe Gln Arg
             165
                               170
Leu Gly Ser Asn Ser Ala Leu Thr Thr Ser Asn Ile Ala Ser Phe Glu
                            185
Glu Ala Phe Ile Cys Leu Gln Lys Leu Met Ala Ala Val Arg Asp Ile
                       200
Leu Glu Gly Ile Gln Arg Ile Leu Ala Pro Asn Ser Asn Tyr Gln Asp
                     215
Val Glu Thr Leu Tyr Asn Phe Leu Ile Lys Tyr Glu Val Asn Lys Asn
   230
                        235 240
Val Lys Phe Thr Ala Gln Glu Ile Tyr Asp Cys Val Ser Gln Thr Glu
           245
                               250
Tyr Arg Glu Lys Leu Thr Ile Gly Cys Arg Gln Leu Val Glu Met Glu
Tyr Thr Met Gln Gln Cys Asn Ala Ser Val Tyr Met Glu Ala Lys Asn
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Arg Gly Trp Cys Glu Asp Met Leu Asn Tyr Arg Ile *
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<210> 1323

<211> 85

<212> PRT

<213> Homo sapiens

<400> 1323

<210> 1324

<211> 46

<212> PRT

<213> Homo sapiens

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Met Leu His His Ser Gln Leu Ile Phe Val Phe Leu Val Gln Thr Gly
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Phe His His Val Ala Leu Ser Gly Phe Lys Leu Leu Ala Ser Ser Asn
                                25
Leu Pro Thr Leu Asp Pro Lys Val Leu Gly Leu Gln Val *
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     <211> 87
     <212> PRT
     <213> Homo sapiens
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Met Gly Leu Ser Lys Ala Phe Leu Ile Thr Arg Thr Val Phe Leu Ile
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                                    10
Ser Ser Leu Ser Phe Tyr Ser Phe Leu Gly Phe Pro Ser Leu Cys Phe
                                25
Thr Gly Ser Cys Met Leu Ser Thr Leu Phe Ile Arg Ala Leu Ser Ile
                            40
Leu Val Ile Ile Val Leu Asn Ser Arg Ser Asp Lys Ser Asn Thr Pro
                        55
                                           60
Ala Ile Ser Glu Ser Gly Ser Asp Ala Cys Ser Phe Ser Ser Asn Phe
                                        75
Val Phe Cys Leu Leu Val *
                85 86
    <210> 1326
    <211> 69 ·
    <212> PRT
    <213> Homo sapiens
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Met Ser Leu Phe Leu Phe Leu Met Phe Gln Val Leu Ser Glu Val
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Ser Trp Gly Gly Val Gly Ser Val Ser Asn Gln Gly Leu Glu His His
                                25
Glu Ile Val Thr Pro Asp Leu Gln Ser Leu Ala Gly Gly Trp Thr Gly
                            40
                                               45
Gly Arg Glu Arg Gly Phe Leu Phe Thr Phe Asn Ile Phe Leu Gln Lys
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                        55
Lys Gln Thr Ile *
65
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    <210> 1327
    <211> 103
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<210> 1328 <211> 52 <212> PRT

<213> Homo sapiens

<210> 1329 <211> 204 <212> PRT <213> Homo sapiens

Glu Leu Thr Asn Gln Val Leu Glu Met Arg Gly Thr Ala Ala Gly Met 105 Asp Leu Trp Val Thr Phe Glu Ile Arg Glu His Gly Glu Leu Glu Arg 120 125 Pro Leu His Pro Lys Glu Lys Val Leu Glu Gln Ala Leu Gln Trp Cys 135 Gln Leu Pro Glu Pro Cys Ser Ala Ser Leu Leu Leu Lys Lys Val Pro 150 155 Leu Ala Gln Ala Gly Cys Leu Phe Thr Gly Ile Arg Arg Glu Ser Pro 165 170 Arg Val Gly Leu Phe Ala Val Phe Val Arg Ser His Leu Ala Cys Trp 180 185 Gly Ser Arg Phe Gln Glu Arg Phe Phe Leu Val Ala 195 200

<210> 1330 <211> 199 <212> PRT <213> Homo sapiens

<400> 1330 Met Pro Val Pro Ala Leu Cys Leu Leu Trp Ala Leu Ala Met Val Thr Arg Pro Ala Ser Ala Ala Pro Met Gly Gly Pro Glu Leu Ala Gln His 2.0 Glu Glu Leu Thr Leu Leu Phe His Gly Thr Leu Gln Leu Gly Gln Ala Leu Asn Gly Val Tyr Arg Thr Thr Glu Gly Arg Leu Thr Lys Ala Arg 55 - Asn Ser Leu Gly Leu Tyr Gly Arg Thr Ile Glu Leu Leu Gly Gln Glu 70 Val Ser Arg Gly Arg Asp Ala Ala Gln Glu Leu Arg Ala Ser Leu Leu 90 Glu Thr Gln Met Glu Glu Asp Ile Leu Gln Leu Gln Ala Glu Ala Thr 105 Ala Glu Val Leu Gly Glu Val Ala Gln Ala Gln Lys Val Leu Arg Asp 120 125 Ser Val Gln Arg Leu Glu Val Gln Leu Arg Ser Ala Trp Leu Gly Pro 135 140 Ala Tyr Arg Glu Phe Glu Val Leu Lys Ala His Ala Asp Lys Gln Ser 150 155 His Ile Leu Trp Ala Leu Thr Gly His Val Gln Arg Gln Arg Glu 165 170 Met Val Ala Gln Gln His Arg Leu Arg Gln Ile Gln Glu Arg Leu His 180 185 Thr Ala Ala Leu Pro Ala * 195 198

<210> 1331 <211> 81 <212> PRT <213> Homo sapiens

<210> 1332 <211> 73 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(73)

<223> Xaa = any amino acid or nothing

<210> 1333 <211> 52 <212> PRT <213> Homo sapiens

<210> 1334

<211> 65 <212> PRT <213> Homo sapiens

<210> 1335 <211> 112 <212> PRT

<213> Homo sapiens

<210> 1336 <211> 105 <212> PRT <213> Homo sapiens

<210> 1337 <211> 57 <212> PRT <213> Homo sapiens

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<210> 1338 <211> 59 <212> PRT <213> Homo sapiens

<210> 1339 <211> 50 <212> PRT <213> Homo sapiens

Tyr * 49

<210> 1340

<211> 81

<212> PRT

<213> Homo sapiens

<400> 1340

Met Pro Leu Ala Cys Thr Gly Leu Asn Thr Gln Arg Phe Ser Tyr Leu 1 5 10 15

Arg Asp Leu Phe Leu Pro Trp Gly Leu Cys Ile Leu Tyr Ser Ile Leu 20 25 30

Ser Ala Ile Phe Pro Asp Leu Ser Ser Ser Ala Lys Leu Pro Ser Leu 35 40 - 45

His Ile Ala Phe Phe Thr Leu Phe Lys Val Thr Lys Gly Thr Ser Pro 50 55 60

Lys Ala Thr Asp Val Pro Val Ala Cys Phe Ile Asn His Asn Arg Thr 65 70 75 80

<210> 1341

<211> 60

<212> PRT

<213> Homo sapiens

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Gln Leu Thr Thr Ser Leu Lys Arg Lys Ser Gly Glu Gly Asp Arg Glu 20 25 . 30

Ser Pro Ala Ser Trp Phe Ser Pro Phe Ser Gln Met Phe Phe Leu Ile 35 40 45

Asn Thr Ile Leu Leu Pro Phe Lys Ile Pro Ile * 50 55 59

<210> 1342

<211> 49

<212> PRT

<213> Homo sapiens

<400> 1342

Met Leu Ser Leu Phe Ile Phe Leu Arg Phe Leu Pro Leu Gly Phe Cys

1 10 15

Trp Lys Glu Leu His Pro Glu Ala Glu Gln Ser Glu Lys Val Asp Phe
20 25 30

Arg Lys Pro Trp Tyr Leu Thr Gly His Ala Ala Ser Leu Gly Ala Asp
35 40 45 48

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<210> 1343
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     <213> Homo sapiens
    <400> 1343
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                                    10
Leu Leu Phe Met Ala His Arg Leu Arg Gln Arg Arg Glu Arg Ile
           20
                               25
Glu Ser Leu Ile Gly Ala Asn Leu His His Phe Asn Leu Gly Arg Arg
                           40
Ile Pro Gly Phe Asp Tyr Gly Pro Asp Gly Phe Gly Thr Gly Leu Thr
                        55
Pro Leu Ala Phe Phe
65
                69
    <210> 1344
    <211> 99
    <212> PRT
    <213> Homo sapiens
    <400> 1344
Met Phe Leu Ser Leu Ser Leu Thr Leu Cys Leu Cys Phe Ser Phe Phe
1 5
                                   10
Cys Leu Tyr Leu Ser Leu Ala Leu Tyr Leu Gly Ser Phe Phe Cys Leu
                               25
Pro Phe His Val Ser Val Phe Leu Cys Leu Phe Pro Ser Val Leu Phe
                           40
                                               45
Leu Ser Val Ala Leu Gly Ser Pro Glu Asn His Ile Ser Trp Arg Lys
                       55
Val Gly Glu Glu Leu Lys Leu Ala Ser His Arg Asn Phe Cys Ser Leu
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                                       75
Met Gln Lys Met Arg Ser Asn Lys Pro Ser Pro Ser Arg Pro Arg Gly
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Trp Ala *
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    <210> 1345
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<400> 1345 Lvs Val Leu

<211> 112 <212> PRT

<213> Homo sapiens

Met Lys Val Leu Trp Ala Gly Val Leu Gly Thr Phe Leu Ala Gly Cys

1 10 15

Gln Ala Lys Val Glu Gln Ala Val Glu Thr Glu Pro Glu Pro Glu Leu
20 25 30

 Cys
 Gln
 Gln
 Thr
 Glu
 Trp
 Lys
 Ser
 Gly
 Gln
 Arg
 Trp
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<210> 1346 <211> 360 <212> PRT <213> Homo sapiens

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<210> 1347 <211> 84 <212> PRT

<213> Homo sapiens

<400> 1347 Met Ile Leu Ser Leu Tyr Tyr Lys Leu Phe Gly Lys Leu Ala Val Ala 1 5 10 Thr Ile Glu Ile Leu His Cys Leu Cys Tyr Ile Glu Phe Val Ile Ile 20 25 Phe Lys Gly Phe Lys Lys Ile Pro Ile Cys Phe Phe Ser Phe Leu Phe 45 Ser Phe Val Pro His His Leu Asn Tyr Leu Gly Lys Tyr His Ser Ser 55 Lys Phe Glu Tyr Cys Leu Ser Asn Lys Lys Cys Glu Arg Tyr Glu 70 75 Glu Glu Arg * 83

<210> 1348 <211> 65 <212> PRT <213> Homo sapiens

<210> 1349 <211> 58 <212> PRT <213> Homo sapiens

<210> 1350 <211> 60 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(60)

<223> Xaa = any amino acid or nothing

<210> 1351 <211> 56 <212> PRT <213> Homo sapiens

<210> 1352 <211> 701 <212> PRT <213> Homo sapiens

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PCT/US01/02687 WO 01/54477

Phe Asn Arg Val Ile Leu Ser Met Lys Arg Gly Gln Glu Tyr Thr Asp 470 475 Tyr Ile Asn Ala Ser Phe Ile Asp Gly Tyr Arg Gln Lys Asp Tyr Phe 485 490 Ile Ala Thr Gln Gly Pro Leu Ala His Thr Val Glu Asp Phe Trp Arg 500 505 Met Ile Trp Glu Trp Lys Ser His Thr Ile Val Met Leu Thr Glu Val 520 Gln Glu Arg Glu Gln Asp Lys Cys Tyr Gln Tyr Trp Pro Thr Glu Gly 535 540 Ser Val Thr His Gly Glu Ile Thr Ile Glu Ile Lys Asn Asp Thr Leu 550 555 Ser Glu Ala Ile Ser Ile Arg Asp Phe Leu Val Thr Leu Asn Gln Pro 565 570 Gln Ala Arg Gln Glu Glu Gln Val Arg Val Arg Gln Phe His Phe 585 His Gly Trp Pro Glu Ile Gly Ile Pro Ala Glu Gly Lys Gly Met Ile 600 Asp Leu Ile Ala Ala Val Gln Lys Gln Gln Gln Gln Thr Gly Asn His Pro Ile Thr Val His Cys Ser Ala Gly Ala Gly Arg Thr Gly Thr Phe 630 635 Ile Ala Leu Ser Asn Ile Leu Glu Arg Val Lys Ala Glu Gly Leu Leu 650 Asp Val Phe Gln Ala Val Lys Ser Leu Arg Leu Gln Arg Pro His Met 665 Val Gln Thr Leu Glu Gln Tyr Glu Phe Cys Tyr Lys Val Val Gln Asp 680 Phe Ile Asp Ile Phe Ser Asp Tyr Ala Asn Phe Lys 695

<210> 1353 <211> 49

<212> PRT

<213> Homo sapiens

<400> 1353

Met Ala Phe Leu Tyr His Val Ala Tyr Val Leu Val Cys Met Leu Gly 10 5 Leu Phe Cys His Glu Phe Phe Tyr Ser Phe Leu Leu Phe Glu Ser Val 20 25 Tyr Arg His Gln Thr Leu Leu Asn Asp Ile Pro Cys Val Lys Leu Met 40

<210> 1354

<211> 58

<212> PRT

<213> Homo sapiens

<400> 1354

Met Ser Val Cys Lys Tyr Thr Val Tyr Gly Phe Phe Ile Phe Ala Phe

<210> 1355 <211> 4261 <212> PRT

<213> Homo sapiens

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	70	~ 7	~	-		_	_	_		~7	_		0 3	7	T
ser	Asp	GIY	Cys 340	Lys	Thr	Pro	Lys	Leu 345	Ile	Glu	Lys	Leu	350	Asp	Leu
Asp	Val	Val 355	Lys	Val	Arg	Cys	Gly 360	Ser	Gln	Phe	Ser	Ile 365	Ala	Leu	Thr
Lys	Asp 370	Gly	Gln	Val	Tyr	Ser 375	Trp	Gly	Lys	Gly	Asp 380	Asn	Gln	Arg	Leu
Gly 385		Gly	Thr	Glu	Glu 390		Val	Arg	Tyr	Pro		Leu	Leu	Glu	Gly 400
	Gln	Gly	Lys	Lys 405		Ile	Asp	Val			Gly	Ser	Thr	His	
Leu	Ala	Leu	Thr		Asp	Ser	Glu		410 His	Ser	Trp	Gly			Asp
Gln	Cys		420 His	Phe	Asp	Thr		425 Arg	Val	Thr	Lys		430 Glu	Pro	Ala
77.	T 011	435	<i>α</i> 1	T	7	m}	440	773 -	~7 -	37-3	~ 3	445	70.7 -	0	~ 1
	450		Gly			455	-				460			_	-
Pro 465	Ala	Gln	Ser	Phe	Ala 470	Trp	Ser	Ser	Cys	Ser 475	Glu	Trp	Ser	Ile	Gly 480
Leu	Arg	Val	Pro	Phe 485	Val	Val	Asp	Ile	Cys 490	Ser	Met	Thr	Phe	Glu 495	Gln
Leu	Asp	Leu	Leu 500	Leu	Arg	Gln	Val	Ser 505	Glu	Gly	Met	Asp	Gly 510	Ser	Ala
Asp	Trp	Pro 515	Pro	Pro	Gln	Glu			Cys	Val	Ala			Thr	Leu
Δen	T.e.11		Arg	T.611	Gln	T.e.u	520	λla	בות	Tlo	Sor	525 Trie	Gln	17 - 7	Acn
	530		_			535					540				_
Pro 545	Glu	Phe	Leu	Gly	Leu 550	Gly	Leu	Gly	Ser	Ile 555	Leu	Leu	Asn	Ser	Leu 560
Lys	Gln	Thr	Val	Val 565	Thr	Leu	Ala	Ser	Ser 570	Ala	Gly	Val	Leu	Ser 575	Thr
Val	Gln	Ser	Ala 580		Gln	Ala	Val	Leu 585		Ser	Gly	Trp	Ser 590		Leu
Leu	Pro	Thr 595	Ala	Glu	Glu	Arg	Ala 600		Ala	Leu	Ser	Ala 605		Leu	Pro
Cvs	Ala		Ser	Glv	Asn	Glu		Δsn	Tle	Ser	Pro		Ara	Ara	Phe
-1-	610		201	011		615	vul	11011	110		620				- 110
Met 625	Ile	Asp	Leu	Leu	Val 630	Gly	Ser	Leu	Met	Ala 635	Asp	Gly	Gly	Leu	Glu 640
	Ala	Leu	His	Ala 645		Ile	Thr	Ala	Glu 650		Gln	Asp	Ile	Glu 655	
Lys	Lys		Ala	Gln	_		_		Ile	_				Ala	Asn
Ala	Ser		660 Phe									Lys			Ile
_	1	675		_		_	680		_		_	685	_	_	-
Asn	1'nr 690	GIY	Ile	Cys	GIu	Ser 695	Ser	GГУ	Lys	GIn	Cys 700	Leu	Pro	Leu	Val
Gln 705	Leu	Ile	Gln	Gln	Leu 710	Leu	Arg	Asn	Ile	Ala 715	Ser	Gln	Thr	Val	Ala 720
Arg	Leu	Lys	Asp	Val 725		Arg	Arg	Ile	Ser 730		Cys	Leu	Asp	Phe 735	
Gln	His	Ser	Arg 740		Arg	Ser	Ala	Ser 745		Asp	Trp	Leu	Leu 750		Phe
Gln	Arg	Leu 755	Leu	Ile	Ser	Lys	Leu 760		Pro	Gly	Glu	Ser 765		Gly	Gln
Thr		_	Ile	Ser	Ser			Leu	Met	Gly			Ser	Leu	Leu
	770 Lys	Tyr	Thr	Ala		775 Leu	Cys	Thr	His		780 Gly	Asp	Ile	Leu	
785 Val	ר ד ת	בומ	9 6~	Tla	790	C.~	wh~	Co~	m	795	u:~	Dho	74.7	G] v	800 Val
val	MIA	AIA	Ser	11e	HIG	Ser.	TILE	ser	тъ	Arg	пта	File	ATA	GIU	val

				805					810					815	
Ala	Tyr	Ile	Val 820	Glu	Gly	Asp	Phe	Thr 825	Gly	Val	Leu	Leu	Pro 830	Glu	Leu
Val	Val	Ser 835	Ile	Val	Leu	Leu	Leu 840	Ser	Lys	Asn	Ala	Asp 845	Leu	Met	Gln
	Ala 850					855					860				-
865	Phe				870		-			875					880
	Ala			885					890					895	-
	Asn		900					905	_	_		_	910		
	Asn	915		-			920					925			-
	Ile 930					935					940				
945	Gln			_	950	_				955					960
	Phe		_	965	•				970					975	
	Leu		980					985					990		
	Ser	995					1000				3	1005			
_	Leu 1010	His	Ala	Ser	_	Leu 1015	Ala	Met	Ser		Pro LO20	Leu	Ser	Pro	Val
Glu 1025	Ile	Glu	Cys		Lys 1030	Trp	Leu	Gln				Phe	Ser		Gly 1040
	Gln	Thr				His	Tyr				Glu	Glu			
Asp	His		Ser 1060	Ser	Pro	Gly	_	Thr 1065	Pro	Ala	Ser	-	Ser 1070	Arg	Leu
Cys	Ser	His LO75	Arg	Arg	Ala		Gly 1080	Asp	His	Ser		Ala 1085	Phe	Leu	Gln
	Ile 1090	Ala	Asp	Asn		Ile 1095	Gln	Asp	His		Val L100	Lys	Asp	Phe	Leu
1105	Gln			1	1110				3	L115				1	120
	Phe]	1125				J	1130				J	1135	
	Cys	1	140				1	L145				1	150		
		155	_			Ī	1160]	165			
1	Lys 1170					175	-	_		1	1180			_	_
1185	Leu			3	1190				1	1195				3	200
	Ala		1	L205				1	1210				1	1215	
	Ala	1	220				1	1225				· 1	230		
		.235				1	L240]	245			
J	Arg 1250		_	-	1	L255	_	_		1	1260				
1265	Lys	тте	чτλ		GIU 1270	GIU	ser	qza		G1u L275	GIU	ATA	cys		Leu 1280

Pro His Ser Pro Ile Asn Val Asp Lys Arg Pro Ile Ala Ile Lys Ser 1285 1290 1295 Pro Lys Asp Lys Trp Gln Pro Leu Ser Thr Val Thr Gly Val His 1300 1305 1310 Lys Tyr Lys Trp Leu Lys Gln Asn Val Gln Gly Leu Tyr Pro Gln Ser 1315 1320 1325 Pro Leu Leu Ser Thr Ile Ala Glu Phe Ala Leu Lys Glu Glu Pro Val 1335 1340 Asp Val Glu Lys Met Arg Lys Cys Leu Leu Lys Gln Leu Glu Arg Ala 1350 1355 1360 Glu Val Arg Leu Glu Gly Ile Asp Thr Ile Leu Lys Leu Ala Ser Lys 1365 1370 Asn Phe Leu Leu Pro Ser Val Gln Tyr Ala Met Phe Cys Gly Trp Gln 1380 1385 1390 Arg Leu Ile Pro Glu Gly Ile Asp Ile Gly Glu Pro Leu Thr Asp Cys 1395 1400 1405 Leu Lys Asp Val Asp Leu Ile Pro Pro Phe Asn Arg Met Leu Leu Glu 1410 1415 1420 Val Thr Phe Gly Lys Leu Tyr Ala Trp Ala Val Gln Asn Ile Arg Asn 1430 1435 Val Leu Met Asp Ala Ser Ala Thr Phe Lys Glu Leu Gly Ile Gln Pro 1445 1450 Val Pro Leu Gln Thr Ile Thr Asn Glu Asn Pro Ser Gly Pro Ser Leu 1465 1470 Gly Thr Ile Pro Gln Ala Arg Phe Leu Leu Val Met Leu Ser Met Leu 1475 1480 1485 Thr Leu Gln His Gly Ala Asn Asn Leu Asp Leu Leu Leu Asn Ser Gly 1495 1500 Met Leu Ala Leu Thr Gln Thr Ala Leu Arg Leu Ile Gly Pro Ser Cys 1510 1515 Asp Asn Val Glu Glu Asp Met Asn Ala Ser Ala Gln Gly Ala Ser Ala 1525 1530 1535 Thr Val Leu Glu Glu Thr Arg Lys Glu Thr Ala Pro Val Gln Leu Pro 1540 1545 1550 Val Ser Gly Pro Glu Leu Ala Ala Met Met Lys Ile Gly Thr Arg Val 1555 1560 1565 Met Arg Gly Val Asp Trp Lys Trp Gly Asp Gln Asp Gly Pro Pro Pro 1570 1575 1580 Gly Leu Gly Arg Val Ile Gly Glu Leu Gly Glu Asp Gly Trp Ile Arg 1590 1595 Val Gln Trp Asp Thr Gly Ser Thr Asn Ser Tyr Arg Met Gly Lys Glu 1605 1610 1615 Gly Lys Tyr Asp Leu Lys Leu Ala Glu Leu Pro Ala Ala Ala Gln Pro 1620 1625 1630 Ser Ala Glu Asp Ser Asp Thr Glu Asp Asp Ser Glu Ala Glu Gln Thr 1635 1640 1645 Glu Arg Asn Ile His Pro Thr Ala Met Met Phe Thr Ser Thr Ile Asn 1650 1655 1660 Leu Leu Gln Thr Leu Cys Leu Ser Ala Gly Val His Ala Glu Ile Met 1670 1675 Gln Ser Glu Ala Thr Lys Thr Leu Cys Gly Leu Leu Arg Met Leu Val 1685 1690 1695 Glu Ser Gly Thr Thr Asp Lys Thr Ser Ser Pro Asn Arg Leu Val Tyr 1700 1705 1710 Arg Glu Gln His Arg Ser Trp Cys Thr Leu Gly Phe Val Arg Ser Ile 1715 1720 1725 Ala Leu Thr Pro Gln Val Cys Gly Ala Leu Ser Ser Pro Gln Trp Ile 1730 1735 1740 Thr Leu Leu Met Lys Val Val Glu Gly His Ala Pro Phe Thr Ala Thr

1750 1755 Ser Leu Gln Arg Gln Ile Leu Ala Val His Leu Leu Gln Ala Val Leu 1765 1770 1775 Pro Ser Trp Asp Lys Thr Glu Arg Ala Arg Asp Met Lys Cys Leu Val 1780 1785 1790 Glu Lys Leu Phe Asp Phe Leu Gly Ser Leu Leu Thr Thr Cys Ser Ser 1795 1800 1805 Asp Val Pro Leu Arg Glu Ser Thr Leu Arg Arg Arg Val Arg 1810 1815 1820 Pro Gln Ala Ser Leu Thr Ala Thr His Ser Ser Thr Leu Ala Glu Glu 1830 1835 1840 Val Val Ala Leu Leu Arg Thr Leu His Ser Leu Thr Gln Trp Asn Gly 1845 1850 1855 Leu Ile Asn Lys Tyr Ile Asn Ser Gln Leu Arg Ser Ile Thr His Ser 1860 1865 1870 Phe Val Gly Arg Pro Ser Glu Gly Ala Gln Leu Glu Asp Tyr Phe Pro 1875 1880 1885 Asp Ser Glu Asn Pro Glu Val Gly Gly Leu Met Ala Val Leu Ala Val 1890 1895 1900 Ile Gly Gly Ile Asp Gly Arg Leu Arg Leu Gly Gly Gln Val Met His 1905 1910 1915 1920 Asp Glu Phe Gly Glu Gly Thr Val Thr Arg Ile Thr Pro Lys Gly Lys 1925 1930 1935 Ile Thr Val Gln Phe Ser Asp Met Arg Thr Cys Arg Val Cys Pro Leu 1940 1945 1950 Asn Gln Leu Lys Pro Leu Pro Ala Val Ala Phe Asn Val Asn Asn Leu 1955 1960 1965 Pro Phe Thr Glu Pro Met Leu Ser Val Trp Ala Gln Leu Val Asn Leu 1970 1975 1980 Ala Gly Ser Lys Leu Glu Lys His Lys Ile Lys Lys Ser Thr Lys Gln 1985 1990 1995 2000 Ala Phe Ala Gly Gln Val Asp Leu Asp Leu Leu Arg Cys Gln Gln Leu 2005 2010 2015 Lys Leu Tyr Ile Leu Lys Ala Gly Arg Ala Leu Leu Ser His Gln Asp 2020 2025 2030 Lys Leu Arg Gln Ile Leu Ser Gln Pro Ala Val Gln Glu Thr Gly Thr 2035 2040 2045 Val His Thr Asp Asp Gly Ala Val Val Ser Pro Asp Leu Gly Asp Met 2050 2055 2060 Ser Pro Glu Gly Pro Gln Pro Pro Met Ile Leu Leu Gln Gln Leu Leu 2070 2075 Ala Ser Ala Thr Gln Pro Ser Pro Val Lys Ala Ile Phe Asp Lys Gln 2085 2090 2095 Glu Leu Glu Ala Ala Ala Leu Ala Val Cys Gln Cys Leu Ala Val Glu 2105 2110 Ser Thr His Pro Ser Ser Pro Gly Phe Glu Asp Cys Ser Ser Ser Glu 2115 2120 2125 Ala Thr Thr Pro Val Ala Val Gln His Ile His Pro Ala Arg Val Lys 2130 2135 2140 Arg Arg Lys Gln Ser Pro Val Pro Ala Leu Pro Ile Val Val Gln Leu 2150 2155 Met Glu Met Gly Phe Ser Arg Arg Asn Ile Glu Phe Ala Leu Lys Ser 2165 2170 2175 Leu Thr Gly Ala Ser Gly Asn Ala Ser Ser Leu Pro Gly Val Glu Ala 2180 2185 2190 Leu Val Gly Trp Leu Leu Asp His Ser Asp Ile Gln Val Thr Glu Leu 2195 2200 2205 Ser Asp Ala Asp Thr Val Ser Asp Glu Tyr Ser Asp Glu Glu Val Val 2215 2220

Glu Asp Val Asp Asp Ala Ala Tyr Ser Met Ser Thr Gly Ala Val Val 2225 2230 2235 2240 Thr Glu Ser Gln Thr Tyr Lys Lys Arg Ala Asp Phe Leu Ser Asn Asp 2245 2250 Asp Tyr Ala Val Tyr Val Arg Glu Asn Ile Gln Val Gly Met Met Val 2260 2265 Arg Cys Cys Arg Ala Tyr Glu Glu Val Cys Glu Gly Asp Val Gly Lys 2275 2280 2285 Val Ile Lys Leu Asp Arg Asp Gly Leu His Asp Leu Asn Val Gln Cys 2295 2300 Asp Trp Gln Gln Lys Gly Gly Thr Tyr Trp Val Arg Tyr Ile His Val 2310 2315 Glu Leu Ile Gly Tyr Pro Pro Pro Ser Ser Ser His Ile Lys Ile 2325 2330 2335 Gly Asp Lys Val Arg Val Lys Ala Ser Val Thr Thr Pro Lys Tyr Lys 2340 2345 2350 Trp Gly Ser Val Thr His Gln Ser Val Gly Val Val Lys Ala Phe Ser 2355 2360 - 2365 Ala Asn Gly Lys Asp Ile Ile Val Asp Phe Pro Gln Gln Ser His Trp 2370 2375 2380 Thr Gly Leu Leu Ser Glu Met Glu Leu Val Pro Ser Ile His Pro Gly 2390 2395 Val Thr Cys Asp Gly Cys Gln Met Phe Pro Ile Asn Gly Ser Arg Phe 2405 2410 2415 Lys Cys Arg Asn Cys Asp Asp Phe Asp Phe Cys Glu Thr Cys Phe Lys 2420 2425 2430 Thr Lys Lys His Asn Thr Arg His Thr Phe Gly Arg Ile Asn Glu Pro 2435 2440 2445 Gly Gln Ser Ala Val Phe Cys Gly Arg Ser Gly Lys Gln Leu Lys Arg 2455 2460 Cys His Ser Ser Gln Pro Gly Met Leu Leu Asp Ser Trp Ser Arg Met 2465 2470 2475 Val Lys Ser Leu Asn Val Ser Ser Ser Val Asn Gln Ala Ser Arg Leu 2485 2490 2495 Ile Asp Gly Ser Glu Pro Cys Trp Gln Ser Ser Gly Ser Gln Gly Lys 2500 2505 2510 His Trp Ile Arg Leu Glu Ile Phe Pro Asp Val Leu Val His Arg Leu 2520 2525 Lys Met Ile Val Asp Pro Ala Asp Ser Ser Tyr Met Pro Ser Leu Val 2535 2540 Val Val Ser Gly Gly Asn Ser Leu Asn Asn Leu Ile Glu Leu Lys Thr 2550 2555 2560 Ile Asn Ile Asn Pro Ser Asp Thr Thr Val Pro Leu Asn Asp Tyr 2565 2570 2575 Thr Glu Tyr His Arg Tyr Ile Glu Ile Ala Ile Lys Gln Cys Arg Ser 2580 2585 2590 Ser Gly Ile Asp Cys Lys Ile His Gly Leu Ile Leu Leu Gly Arg Ile 2600 2605 Arg Ala Glu Glu Asp Leu Ala Ala Val Pro Phe Leu Ala Ser Asp 2615 2620 Asn Glu Glu Glu Asp Glu Lys Gly Asn Ser Gly Ser Leu Ile Arg 2630 2635 Lys Lys Ala Ala Gly Leu Glu Ser Ala Ala Thr Ile Arg Thr Lys Val 2645 2650 Phe Val Trp Gly Leu Asn Asp Lys Asp Gln Leu Gly Gly Leu Lys Gly 2660 2665 Ser Lys Ile Lys Val Pro Ser Phe Ser Glu Thr Leu Ser Ala Leu Asn 2680 2685 Val Val Gln Val Ala Gly Gly Ser Lys Ser Leu Phe Ala Val Thr Val

2690 2695 2700 Glu Gly Lys Val Tyr Ala Cys Gly Glu Ala Thr Asn Gly Arg Leu Gly **2705 2710 2715 2720** Leu Gly Ile Ser Ser Gly Thr Val Pro Ile Pro Arg Gln Ile Thr Ala 2725 2730 2735 Leu Ser Ser Tyr Val Val Lys Lys Val Ala Val His Ser Gly Gly Arg 2740 2745 2750 His Ala Thr Ala Leu Thr Val Asp Gly Lys Val Phe Ser Trp Gly Glu 2755 2760 2765 Gly Asp Asp Gly Lys Leu Gly His Phe Ser Arg Met Asn Cys Asp Lys 2770 2775 2780 Pro Arg Leu Ile Glu Ala Leu Lys Thr Lys Arg Ile Arg Asp Ile Ala 2785 2790 2795 2800 Cys Gly Ser Ser His Ser Ala Ala Leu Thr Ser Ser Gly Glu Leu Tyr 2805 2810 2815 Thr Trp Gly Leu Gly Glu Tyr Gly Arg Leu Gly His Gly Asp Asn Thr 2825 Thr Gln Leu Lys Pro Lys Met Val Lys Val Leu Leu Gly His Arg Val 2835 2840 2845 Ile Gln Val Ala Cys Gly Ser Arg Asp Ala Gln Thr Leu Ala Leu Thr 2855 2860 Asp Glu Gly Leu Val Phe Ser Trp Gly Asp Gly Asp Phe Gly Lys Leu 2870 2875 2880 Gly Arg Gly Gly Ser Glu Gly Cys Asn Ile Pro Gln Asn Ile Glu Arg 2885 2890 2895 Leu Asn Gly Gln Gly Val Cys Gln Ile Glu Cys Gly Ala Gln Phe Ser 2900 2905 2910 Leu Ala Leu Thr Lys Ser Gly Val Val Trp Thr Trp Gly Lys Gly Asp 2915 2920 2925 Tyr Phe Arg Leu Gly His Gly Ser Asp Val His Val Arg Lys Pro Gln 2930 2935 2940 Val Val Glu Gly Leu Arg Gly Lys Lys Ile Val His Val Ala Val Gly 2950 2955 2960 Ala Leu His Cys Leu Ala Val Thr Asp Ser Gly Gln Val Tyr Ala Trp 2965 2970 2975 Gly Asp Asn Asp His Gly Gln Gln Gly Asn Gly Thr Thr Thr Val Asn 2980 2985 2990 Arg Lys Pro Thr Leu Val Gln Gly Leu Glu Gly Gln Lys Ile Thr Arg 2995 3000 3005 Val Ala Cys Gly Ser Ser His Ser Val Ala Trp Thr Thr Val Asp Val 3010 3015 3020 Ala Thr Pro Ser Val His Glu Pro Val Leu Phe Gln Thr Ala Arg Asp 3025 3030 3035 3040 Pro Leu Gly Ala Ser Tyr Leu Gly Val Pro Ser Asp Ala Asp Ser Ser 3045 3050 3055 Ala Ser Asn Lys Ile Ser Gly Ala Ser Asn Ser Lys Pro Asn Arg 3060 3065 3070 Pro Ser Leu Ala Lys Ile Leu Leu Ser Leu Asp Gly Asn Leu Ala Lys 3080 3085 Gln Gln Ala Leu Ser His Ile Leu Thr Ala Leu Gln Ile Met Tyr Ala 3095 3100 Arg Asp Ala Val Val Gly Ala Leu Met Pro Ala Ala Met Ile Ala Pro 3110 3115 3120 Val Glu Cys Pro Ser Phe Ser Ser Ala Ala Pro Ser Asp Ala Ser Ala 3125 3130 3135 Met Ala Ser Pro Met Asn Gly Glu Cys Met Leu Ala Val Asp Ile 3140 3145 3150 Glu Asp Arg Leu Ser Pro Asn Pro Trp Gln Glu Lys Arg Glu Ile Val 3155 3160 3165

Ser Ser Glu Asp Ala Val Thr Pro Ser Ala Val Thr Pro Ser Ala Pro 3170 3175 3180 Ser Ala Ser Ala Arg Pro Phe Ile Pro Val Thr Asp Asp Leu Gly Ala 3190 3195 Ala Ser Ile Ile Ala Glu Thr Met Thr Lys Thr Lys Glu Asp Val Glu 3205 3210 3215 Ser Gln Asn Lys Ala Ala Gly Pro Glu Pro Gln Ala Leu Asp Glu Phe 3225 Thr Ser Leu Leu Ile Ala Asp Asp Thr Arg Val Val Asp Leu Leu 3235 3240 3245 Lys Leu Ser Val Cys Ser Arg Ala Gly Asp Arg Gly Arg Asp Val Leu 3255 3260 Ser Ala Val Leu Ser Gly Met Gly Thr Ala Tyr Pro Gln Val Ala Asp 3270 3275 Met Leu Glu Leu Cys Val Thr Glu Leu Glu Asp Val Ala Thr Asp 3285 3290 Ser Gln Ser Gly Arg Leu Ser Ser Gln Pro Val Val Glu Ser Ser 3300 3305 His Pro Tyr Thr Asp Asp Thr Ser Thr Ser Gly Thr Val Lys Ile Pro 3320 ' 3315 3325 Gly Ala Glu Gly Leu Arg Val Glu Phe Asp Arg Gln Cys Ser Thr Glu 3335 3340 Arg Arg His Asp Pro Leu Thr Val Met Asp Gly Val Asn Arg Ile Val 3355 3360 3350 Ser Val Arg Ser Gly Arg Glu Trp Ser Asp Trp Ser Ser Glu Leu Arg 3365 3370 3375 Ile Pro Gly Asp Glu Leu Lys Trp Lys Phe Ile Ser Asp Gly Ser Val 3385 3390 Asn Gly Trp Gly Trp Arg Phe Thr Val Tyr Pro Ile Met Pro Ala Ala 3400 3405 Gly Pro Lys Glu Leu Leu Ser Asp Arg Cys Val Leu Ser Cys Pro Ser 3415 3420 Met Asp Leu Val Thr Cys Leu Leu Asp Phe Arg Leu Asn Leu Ala Ser 3430 3435 3440 Asn Arg Ser Ile Val Pro Arg Leu Ala Ala Ser Leu Ala Ala Cys Ala 3445 3450 3455 Gln Leu Ser Ala Leu Ala Ala Ser His Arg Met Trp Ala Leu Gln Arg 3460 3465 3470 Leu Arg Lys Leu Leu Thr Thr Glu Phe Gly Gln Ser Ile Asn Ile Asn 3475 3480 3485 Arg Leu Leu Gly Glu Asn Asp Gly Glu Thr Arg Ala Leu Ser Phe Thr 3495 3500 Gly Ser Ala Leu Ala Ala Leu Val Lys Gly Leu Pro Glu Ala Leu Gln 3510 3515 Arg Gln Phe Glu Tyr Glu Asp Pro Ile Val Arg Gly Gly Lys Gln Leu 3525 3530 Leu His Ser Pro Phe Phe Lys Val Leu Val Ala Leu Ala Cys Asp Leu 3540 3545 3550 Glu Leu Asp Thr Leu Pro Cys Cys Ala Glu Thr His Lys Trp Ala Trp 3560 3565 Phe Arg Arg Tyr Cys Met Ala Ser Arg Val Ala Val Ala Leu Asp Lys 3575 3580 Arg Thr Pro Leu Pro Arg Leu Phe Leu Asp Glu Val Ala Lys Lys Ile 3590 3595 Arg Glu Leu Met Ala Asp Ser Glu Asn Met Asp Val Leu His Glu Ser 3605 3610 His Asp Ile Phe Lys Arg Glu Gln Asp Glu Gln Leu Val Gln Trp Met 3620 3625 3630 Asn Arg Arg Pro Asp Asp Trp Thr Leu Ser Ala Gly Gly Ser Gly Thr

3635 3640 3645 Ile Tyr Gly Trp Gly His Asn His Arg Gly Gln Leu Gly Gly Ile Glu 3650 3655 3660 Gly Ala Lys Val Lys Val Pro Thr Pro Cys Glu Ala Leu Ala Thr Leu 3665 3670 3675 3680 Arg Pro Val Gln Leu Ile Gly Gly Glu Gln Thr Leu Phe Ala Val Thr 3685 3690 3695 Ala Asp Gly Lys Leu Tyr Ala Thr Gly Tyr Gly Ala Gly Gly Arg Leu 3700 3705 3710 Gly Ile Gly Gly Thr Glu Ser Val Ser Thr Pro Thr Leu Leu Glu Ser 3715 3720 3725 Ile Gln His Val Phe Ile Lys Lys Val Ala Val Asn Ser Gly Gly Lys 3730 3735 3740 His Cys Leu Ala Leu Ser Ser Glu Gly Glu Val Tyr Ser Trp Gly Glu **3745 3750 3755 3760** Ala Glu Asp Gly Lys Leu Gly His Gly Asn Arg Ser Pro Cys Asp Arg 3765 3770 3775 Pro Arg Val Ile Glu Ser Leu Arg Gly Ile Glu Val Val Asp Val Ala 3780 3785 3790 Ala Gly Gly Ala His Ser Ala Cys Val Thr Ala Ala Gly Asp Leu Tyr 3795 3800 3805 Thr Trp Gly Lys Gly Arg Tyr Gly Arg Leu Gly His Ser Asp Ser Glu 3810 3815 3820 Asp Gln Leu Lys Pro Lys Leu Val Glu Ala Leu Gln Gly His Arg Val 3825 3830 3835 3840 Val Asp Ile Ala Cys Gly Ser Gly Asp Ala Gln Thr Leu Cys Leu Thr 3845 3850 3855 Asp Asp Asp Thr Val Trp Ser Trp Gly Asp Gly Asp Tyr Gly Lys Leu 3860 3865 3870 Gly Arg Gly Ser Asp Gly Cys Lys Val Pro Met Lys Ile Asp Ser 3875 3880 3885 Leu Thr Gly Leu Gly Val Val Lys Val Glu Cys Gly Ser Gln Phe Ser 3890 3895 3900 Val Ala Leu Thr Lys Ser Gly Ala Val Tyr Thr Trp Gly Lys Gly Asp 3905 3910 3915 Tyr His Arg Leu Gly His Gly Ser Asp Asp His Val Arg Arg Pro Arg 3925 3930 3935 Gln Val Gln Gly Leu Gln Gly Lys Lys Val Ile Ala Ile Ala Thr Gly 3940 3945 3950 Ser Leu His Cys Val Cys Cys Thr Glu Asp Gly Glu Val Tyr Thr Trp 3955 3960 3965 Gly Asp Asn Asp Glu Gly Gln Leu Gly Asp Gly Thr Thr Asn Ala Ile 3970 3975 3980 Gln Arg Pro Arg Leu Val Ala Ala Leu Gln Gly Lys Lys Val Asn Arg 3990 3995 4000 Val Ala Cys Gly Ser Ala His Thr Leu Ala Trp Ser Thr Ser Lys Pro 4005 4010 4015 Ala Ser Ala Gly Lys Leu Pro Ala Gln Val Pro Met Glu Tyr Asn His 4020 4025 4030 Leu Gln Glu Ile Pro Ile Ile Ala Leu Arg Asn Arg Leu Leu Leu Leu 4035 4040 4045 His His Leu Ser Glu Leu Phe Cys Pro Cys Ile Pro Met Phe Asp Leu 4055 4060 Glu Gly Ser Leu Asp Glu Thr Gly Leu Gly Pro Ser Val Gly Phe Asp 4075 Thr Leu Arg Gly Ile Leu Ile Ser Gln Gly Lys Glu Ala Ala Phe Arg 4090 Lys Val Val Gln Ala Thr Met Val Arg Asp Arg Gln His Gly Pro Val 4105

Val Glu Leu Asn Arg Ile Gln Val Lys Arg Ser Arg Ser Lys Gly Gly 4115 4120 4125 Leu Ala Gly Pro Asp Gly Thr Lys Ser Val Phe Gly Gln Met Cys Ala 4135 4140 Lys Met Ser Ser Phe Gly Pro Asp Ser Leu Leu Leu Pro His Arg Val 4150 4155 4160 Trp Lys Val Lys Phe Val Gly Glu Ser Val Asp Asp Cys Gly Gly Gly 4165 4170 4175 Tyr Ser Glu Ser Ile Ala Glu Ile Cys Glu Glu Leu Gln Asn Gly Leu 4180 4185 4190 Thr Pro Leu Leu Ile Val Thr Pro Asn Gly Arg Asp Glu Ser Gly Ala 4200 4205 Asn Arg Asp Cys Tyr Leu Leu Ser Pro Ala Ala Arg Ala Pro Val His 4215 4220 Ser Ser Met Phe Arg Phe Leu Gly Val Leu Gly Ile Ala Ile Arg 4230 4235 Thr Gly Ser Pro Leu Ser Leu Asn Pro Cys Arg Ala Leu Ser Gly Ser 4245 4250 Ser Trp Leu Gly * 4260

<210> 1356

<211> 64

<212> PRT

<213> Homo sapiens

<400> 1356

 Met
 Ser
 Lys
 Val
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 His
 Gly
 Ala
 Pro
 Ala
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 Leu
 Leu
 Val

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 Ser
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 Cys
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 Gly
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 Ile
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 His

 Val
 Thr
 Tyr
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 His
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<223> Xaa = any amino acid or nothing

<400> 1357

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 1
 5
 10
 15

 Leu Ala Thr Leu Lys Val Leu Ser Leu Leu Trp Leu Leu Tyr Tyr Val
 20
 25
 30

 Ala Ser Thr Thr Arg Gln Pro His Ala Val Leu Tyr Gln Asp Pro His
 45

 Ala Gly Pro Leu Trp Val Arg Ser Ser Leu Val Leu Phe Gly Ser Cys

Thr Phe Cys Leu Asn Ile Phe Arg Val Gly Tyr Asp Val Ser His Ile 65 70 75 80

Arg Cys Lys Ser Gln Leu Asp Leu Val Phe Pro Val Ile Glu Met Val 85 90 95

Phe Ile Gly Val Gln Thr Cys Val Leu Trp Lys His Cys Arg Xaa 100 110 111

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<210> 1359 <211> 73 <212> PRT <213> Homo sapiens

<210> 1360 <211> 57 <212> PRT <213> Homo sapiens

Phe Phe Phe Ala Phe Phe Arg Thr * 50 55 56

<210> 1361

<211> 77

<212> PRT

<213> Homo sapiens

<400> 1361

Met Phe Val Leu Phe Leu Ile Leu Val Leu Arg Asn His Phe Leu Val 1 5 10 15

Thr Ile Lys Tyr Gly Val Gly Cys Gly Phe Ile Ile Ser Val Cys Leu 20 25 30

Arg Ala Lys His Phe Asn Phe Asp Glu Ala Gln Phe Val Ser Phe Phe 35 40 45

Leu Cys Asp Ser Cys Phe Cys Leu Leu Arg Asn Leu Pro Thr Gln Arg 50 55 60

Leu Gln Arg Phe Phe Phe Cys Trp Phe Phe Leu Ile * 65 70 75 76

<210> 1362

<211> 106

<212> PRT

<213> Homo sapiens

<400> 1362

Met Gln Asn Arg Thr Gly Leu Ile Leu Cys Ala Leu Ala Leu Leu Met 1 5 10 15

Gly Phe Leu Met Val Cys Leu Gly Ala Phe Phe Ile Ser Trp Gly Ser
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Ile Phe Asp Cys Gln Gly Ser Leu Ile Ala Ala Tyr Leu Leu Pro
35 40 45

Leu Gly Phe Val Ile Leu Leu Ser Gly Ile Phe Trp Ser Asn Tyr Arg
50 55 60

Gln Val Thr Glu Ser Lys Gly Val Leu Arg His Met Leu Arg Gln His 65 70 75 80

Leu Ala His Gly Ala Leu Pro Val Ala Thr Val Asp Arg Ala Ala Leu 85 90 95

Leu Lys Ile Met Cys Lys Gln Leu Leu * 100 105

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<212> PRT

<213> Homo sapiens

<400> 1363

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Leu Ala Thr Leu Cys Ile Asp Arg Gln Val Ser Ser Ser Leu Val

20 25 30

Gln Glu Gly Phe His Ser Lys Ser Cys His Cys Leu Gly Asp Ser Phe
35 40 45

Arg Glu Lys Asn Gln Val Val Gly *
50 55 56

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<213> Homo sapiens

<210> 1365 <211> 58 <212> PRT <213> Homo sapiens

<210> 1366 <211> 58 <212> PRT <213> Homo sapiens

Leu Asp Leu Tyr Ser Ser Leu Phe Phe * 50 55 57

<210> 1367

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1367

35 40 45 47

<210> 1368

<211> 96

<212> PRT

<213> Homo sapiens

<400> 1368

 Met
 Gly
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 Lys
 Ser
 Phe
 Phe
 Phe
 Leu
 Phe
 Leu
 Glu
 Cys
 Arg
 Glu

 Lys
 Gly
 Leu
 His
 Ile
 Pro
 Leu
 Cys
 Thr
 Cys
 Ser
 His
 Ala
 Pro
 Arg
 Pro

 Pro
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 Ala
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 Ser
 Ala
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 Pro
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 Thr
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 Arg
 Gly
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 Leu
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<210> 1369

<211> 76

<212> PRT

<213> Homo sapiens

<400> 1369

 Met
 Trp
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 Val
 Leu
 Ser
 Arg
 Val
 Leu
 Lys
 Asp
 His
 Met
 Ala
 Ser
 Asn
 Ala
 Tyr
 Lys

 Phe
 His
 Ser
 Arg
 Val
 Leu
 Lys
 Asp
 His
 Met
 Ala
 Ser
 Asn
 Ala
 Tyr
 Lys

 Ser
 Ala
 Leu
 Phe
 Phe
 Thr
 Val
 Arg
 Tyr
 Leu
 Glu
 Thr
 Lys
 Gln
 Phe
 Leu

 Arg
 Cys
 Cys
 Cys
 Trp
 Pro
 Asp
 Ala
 Val
 Ala
 His
 Ala
 Cys
 Asn
 Thr

 Leu
 Arg
 Cys
 Cys
 Trp
 Pro
 Asp
 Ala
 Val
 Ala
 His
 Ala
 Cys
 Asn
 Thr

 Leu
 Arg
 Gly
 Gln
 Gly
 Arg
 His
 Ile
 Thr
 *

65 70 75

<210> 1370 <211> 79 <212> PRT

<213> Homo sapiens

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75

<210> 1371 <211> 227 <212> PRT <213> Homo sapiens <221> misc feature <222> (1)...(227) <223> Xaa = any amino acid or nothing

<400> 1371 Met Leu Tyr Phe Gln Leu Val Ile Met Ala Gly Thr Val Leu Leu Ala 5 10 Tyr Tyr Phe Glu Cys Thr Asp Thr Phe Gln Val His Ile Gln Gly Phe 25 Phe Cys Gln Asp Gly Asp Leu Met Lys Pro Tyr Pro Gly Thr Glu Glu 40 45 Glu Ser Phe Ile Thr Pro Leu Val Leu Tyr Cys Val Leu Ala Ala Thr 55 Pro Thr Ala Ile Ile Phe Ile Gly Glu Ile Ser Met Tyr Phe Ile Lys 70 75 Ser Thr Arg Glu Ser Leu Ile Ala Gln Glu Lys Thr Ile Leu Thr Gly 85 90 Glu Cys Cys Tyr Leu Asn Pro Leu Leu Arg Arg Ile Ile Arg Phe Thr 105 100 110 Gly Val Phe Ala Phe Gly Leu Phe Ala Thr Asp Ile Phe Val Asn Ala 120 125 Gly Gln Val Val Thr Gly His Leu Thr Pro Tyr Phe Leu Thr Val Cys 135 Lys Pro Asn Tyr Thr Ser Ala Asp Cys Gln Ala His His Gln Phe Ile 155 Asn Asn Gly Asn Ile Cys Thr Gly Asp Leu Gly Ser Asp Arg Lys Gly 170 Ser Glu Ile Leu Ser Leu Gln Thr Arg Cys Ser Glu His Leu Leu Arg 185

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Leu Ile Trp Pro Arg Cys Ile Phe Thr Arg His Asn Gln Gly Arg Gly
    195
                           200
Gly Ser Ser Met Gly Pro Ser Arg Trp Leu Cys Leu Gly Thr Phe Leu
                      215
His Xaa Leu
225 227
     <210> 1372
     <211> 99
     <212> PRT
     <213> Homo sapiens
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Met Phe Leu Ser Leu Ser Leu Thr Leu Cys Leu Cys Phe Ser Phe Phe
                5
Cys Leu Tyr Leu Ser Leu Ser Leu Tyr Leu Arg Ser Phe Phe Cys Leu
            20
                                25
Pro Phe His Val Ser Val Phe Leu Cys Leu Phe Pro Ser Val Leu Phe
Leu Ser Val Ala Leu Gly Ser Pro Glu Asn His Ile Ser Trp Arg Lys
Val Gly Glu Leu Lys Leu Ala Ser His Arg Asn Phe Cys Ser Leu
                     70
                                        75
Ile Gln Met Met Arg Ser Asn Lys Pro Ser Pro Ser Arg Gln Arg Gly
                 85
                                    90
Trp Ala *
    98
     <210> 1373
     <211> 69
     <212> PRT
     <213> Homo sapiens
     <400> 1373
Met Leu His Thr Pro Gln Thr Cys Arg Pro Gly Leu Cys Val Leu Ala
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Ser Arg Pro Val Leu Tyr Thr Leu Cys Leu Leu Ile Pro Val Leu Cys
           20
                               25
Gly Asp Thr Phe Trp Ala Ser Trp Ser Leu Leu Thr Lys Ala Thr Pro
                            40
Ser Ser Leu Leu Cys Leu Ser Asp Lys Ser Ile Pro Ser Leu Ile Ser
                        55
Lys Gly Asp Ser *
           68
    <210> 1374
    <211> 296
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<212> PRT

<213> Homo sapiens

<400> 1374 Met Arg Ser Lys Ile Met Ile His Ile His Ile Phe Leu Leu Ala Ser - 5 10 Phe Arq Phe Lys Glu His Val Gln Asn Asn Leu Pro Arg Asp Leu Leu 25 Thr Gly Glu Gln Phe Ile Gln Leu Arg Arg Glu Leu Ala Ser Val Asn 40 Gly His Ser Gly Asp Asp Gly Pro Pro Gly Asp Asp Leu Pro Ser Gly 55 Ile Glu Asp Ile Thr Asp Pro Ala Lys Leu Ile Thr Glu Ile Glu Asn 70 75 Met Arg His Arg Ile Ile Glu Ile His Gln Glu Met Phe Asn Tyr Asn 90 85 Glu His Glu Val Ser Lys Arg Trp Thr Phe Glu Glu Gly Ile Lys Arg 105 100 Pro Tyr Phe His Val Lys Pro Leu Glu Lys Ala Gln Leu Lys Asn Trp 120 125 Lys Glu Tyr Leu Glu Phe Glu Ile Glu Asn Gly Thr His Glu Arg Val 135 140 Val Val Leu Phe Glu Arg Cys Val Ile Ser Cys Ala Leu Tyr Glu Glu 150 155 Phe Trp Ile Lys Tyr Ala Lys Tyr Met Glu Asn His Ser Ile Glu Gly 170 Val Arg His Val Phe Ser Arg Ala Cys Thr Ile His Leu Pro Lys Lys 185 Pro Met Val His Met Leu Trp Ala Ala Phe Glu Glu Gln Gln Gly Asn 200 Ile Asn Glu Ala Arg Asn Ile Leu Lys Thr Phe Glu Glu Cys Val Leu 215 220 Gly Leu Ala Met Val Arg Leu Arg Arg Val Ser Leu Glu Arg Arg His 230 235 Gly Asn Leu Glu Glu Ala Glu His Leu Leu Gln Asp Ala Ile Lys Asn 245 250 Ala Lys Ser Asn Asn Glu Ser Ser Phe Tyr Ala Val Lys Leu Ala Arg 265 270 His Leu Phe Lys Ile Gln Lys Asn Leu Pro Lys Ser Arg Lys Val Leu 280 Leu Glu Ala Ile Glu Arg Asp Lys 295 296

<210> 1375 <211> 75 <212> PRT <213> Homo sapiens

<210> 1376 <211> 61 <212> PRT <213> Homo sapiens

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<210> 1377 <211> 110 <212> PRT <213> Homo sapiens

| Met | Trp | Val | Trp | Val | Thr | Ala | Ala | His | Leu | Leu | Cys | Ser | Leu | Ala | Ala | Ala | Ala | Leu | Leu | Cys | Ser | Leu | Ala | Ala | Ala | Ala | Leu | Leu | Cys | Ser | Leu | Ala | Ala | Ala | Ala | Ala | Leu | Leu | Cys | Leu | Arg | Val | Asp | Val | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys

<210> 1378 <211> 47 <212> PRT <213> Homo sapiens

<210> 1379 <211> 140 <212> PRT <213> Homo sapiens

<400> 1379 Met Arg His Pro Ser Pro Trp Pro Phe Leu Phe Phe Cys Phe Val Pro 10 Ala Thr Leu Arg Ser Phe Pro Ser Gly Leu Val Trp Pro Gly Cys Trp 20 25 Trp Glu Pro Arg Ala Ser Pro Ser Ser Leu Ala Pro Gly Met Lys Ser 40 Gln Leu Trp Ala Ala Ala Trp Arg Pro Gly Thr Ser Leu Gln Gly Met 55 Ala Gly Ile Leu Arg Gln Ala Ala Glu Ala Gly Pro Ala Gly Val Ala 75 Leu Ile Leu Ile Lys Gly Thr Gly Asn Glu Glu Pro Leu Gly Pro Leu Pro Ser Arg Cys Leu Cys Pro Pro Pro Glu Glu Pro Arg Phe His Trp 105 Ala Leu Gly Lys Glu Pro Thr Gly Pro Gly Arg Pro Gln Pro Val Gln 120 His His Ile Glu Gly Pro His Pro Val Gly Phe Gly

<210> 1380 <211> 50 <212> PRT <213> Homo sapiens

<210> 1381 <211> 78 <212> PRT <213> Homo sapiens

PCT/US01/02687 WO 01/54477

Val Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 40 Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Asn Pro Ser Gly Pro 55 60 Tyr Gln Lys Lys Pro Val His Glu Lys Lys Glu Val Leu * 70

<210> 1382 <211> 57 <212> PRT <213> Homo sapiens

<400> 1382 Met Leu Thr Thr Leu Leu Leu Leu His Lys Arg Ile Phe Arg Gly 5 10 Asn Phe His Ile Leu His Phe His Ile Cys Ile Gln Ile Lys Lys Gln 2.0 25 30 Ile Pro Ile Leu Glu Asn Asp Leu Phe Lys Met Tyr Thr Val Ser Asn 3.5 40 Lys Ala Lys Thr Arg Thr Trp Ser *

55 56

<210> 1383 <211> 64 <212> PRT <213> Homo sapiens

<400> 1383 Met Val Cys Arg Leu Pro Cys Thr Leu Leu Pro Trp Pro Leu Lys His 5 10 Lys Gln Gly Ala Leu Leu Tyr Ile Cys Pro Ala Ser Leu Pro Ala Phe 25 Asn Pro Arg Asn Leu Ser Val Tyr Leu Leu Phe Ser Ala Ser Glu Ser 40 45 Leu Pro Leu Lys Ser Glu Gln Ala Arg Pro Gly Gly Ser Arg Leu * 50

<210> 1384 <211> 67 <212> PRT <213> Homo sapiens

<400> 1384 Met Leu Ser Phe Val Pro Leu Leu Ser Ser Trp Leu Gly Thr Trp Ile 5 10 Thr Asp Arg Gly Ala Ala Gly Ser Cys Gln Ala Glu Ala Pro Arg Leu 25 20 Ala Gly Glu Thr Ala Gly Gln Arg Val Trp Glu Arg Gly Met Gln Arg Ala Ala Val Gly Lys Ile Leu Asp Pro Lys Gly His Thr Ala Ser

50 55 60 Pro His * 65 66

> <210> 1385 <211> 50 <212> PRT

<213> Homo sapiens

<210> 1386 <211> 123 <212> PRT <213> Homo sapiens

<400> 1386 Met Lys Trp Val Thr Phe Ile Ser Leu Leu Phe Leu Phe Ser Ser Ala 5 10 Tyr Ser Arg Gly Pro Lys Ala Glu Phe Ala Glu Val Ser Lys Leu Val 20 25 Thr Asp Leu Thr Lys Val His Thr Glu Cys Cys His Gly Asp Leu Leu 45 40 Glu Cys Ala Asp Asp Arg Ala Asp Leu Ala Lys Tyr Ile Cys Glu Asn 55 60 Gln Asp Ser Ile Ser Ser Lys Leu Lys Glu Cys Cys Glu Lys Pro Leu 70 75 Leu Glu Lys Ser His Cys Ile Ala Glu Val Glu Asn Asp Glu Met Pro 85 90 Ala Asp Leu Pro Ser Leu Ala Ala Asp Phe Val Glu Ser Lys Asp Val 105 Cys Lys Asn Tyr Ala Glu Ala Lys Asp Val Phe

120

<210> 1387 <211> 65 <212> PRT <213> Homo sapiens

<210> 1388

<211> 56

<212> PRT

<213> Homo sapiens

<400> 1388

<210> 1389

<211> 76

<212> PRT

<213> Homo sapiens

<400> 1389

<210> 1390

<211> 149

<212> PRT

<213> Homo sapiens

<400> 1390

Met Ala Ala Ser Pro Ala Arg Pro Ala Val Leu Ala Leu Thr Gly Leu 1 5 10 15
Ala Leu Leu Leu Leu Cys Trp Gly Pro Gly Gly Ile Ser Gly Asn

20 25 Lys Leu Lys Leu Met Leu Gln Lys Arg Glu Ala Pro Val Pro Thr Lys 40 Thr Lys Val Ala Val Asp Glu Asn Lys Ala Lys Glu Phe Leu Gly Ser 55 Leu Lys Arg Gln Lys Arg Gln Leu Trp Asp Arg Thr Arg Pro Glu Val 75 Gln Gln Trp Tyr Gln Gln Phe Leu Tyr Met Gly Phe Asp Glu Ala Lys 85 90 Phe Glu Asp Asp Ile Thr Tyr Trp Leu Asn Arg Asp Arg Asn Gly His 100 105 Glu Tyr Tyr Gly Asp Tyr Tyr Gln Arg His Tyr Asp Glu Asp Ser Ala 120 Ile Gly Pro Arg Ser Pro Tyr Gly Phe Arg His Gly Ala Ser Val Asn 135 Tyr Asp Asp Tyr * 145 148

<210> 1391 <211> 125 <212> PRT <213> Homo sapiens

<400> 1391 Met Val Met Gly Trp His Trp Pro Gln Gly Leu Gly Leu Ser Leu Ser 10 Leu Cys Pro Ser Asp Leu Asp Gly Trp Val Ser Arg Glu Val Pro Leu 20 25 Leu Asp Arg Pro Gln Ala Leu Pro Pro Cys Val Gln Ile Leu Ser Ala 35 40 Pro Ala Ser Thr Ser Cys Pro Ser Ala Leu Ser Pro Trp His Asp Pro 55 . 60 Gly Leu Pro Val Thr Ser Gln Asn His Phe Ala Trp Phe Pro Leu Gly 70 75 Ser Lys Ala Cys Leu Gly Pro Ser Ile Asp Arg Glu Ala Val Lys Glu 85 90 Ile Asn Ala Glu Glu Gly Val Arg Arg Gln Thr Gln Gly Pro Ile Lys 105 Val Arg Lys Gln Ala Gly Cys Gly Gly Ser Cys Leu * 120

<210> 1392 <211> 56 <212> PRT <213> Homo sapiens

Ile Ile Leu Pro Leu His Pro * 50 55

<210> 1393

<211> 55

<212> PRT

<213> Homo sapiens

<400> 1393

Met Glu Ala Trp Lys Ala Leu Ile Gly Leu Phe Pro Leu Arg Ser Ser 1 10 15

Ala Ser Pro Phe Thr Tyr His Cys Trp Glu Pro Ala Gln Pro Ala His 20 25 30

Gln Glu Phe His Ser Thr Ile Ala Leu Arg Gly Arg Gly Gly Lys Pro

Gln Glu Phe His Ser Thr Ile Ala Leu Arg Gly Arg Gly Gly Lys Pro 35 40 45

Gln Glu Glu Ser Ser Pro * 50 54

<210> 1394

<211> 51

<212> PRT

<213> Homo sapiens

<400> 1394

<210> 1395

<211> 105

<212> PRT

<213> Homo sapiens

<400> 1395

 Met
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Phe Gly Leu Leu Ser Leu Pro Ser Ile
100 105

<210> 1396 <211> 49 <212> PRT

<213> Homo sapiens

<210> 1397 <211> 104 <212> PRT <213> Homo sapiens

<400> 1397 Met Leu Ser Trp Val Phe Pro Gly Ser Val Phe Gly Leu Cys Leu Ser 1 5 10 Val Trp Val Phe Trp His Gln Ala Ser Leu Gly Arg Ala Ser Gly Cys 25 Ala Pro Ala Leu Arg Val Gly Leu Ile Pro Gly Cys Arg Gly Leu Arg 40 45 Ala Glu Leu Phe His Leu Glu Asp Lys Asp Gly Ser Ser Gly Leu Gly 60 Gly Gly Gly Ala Gly His Asp Leu Ile Leu Arg Arg Ala Trp Cys 70 75 Trp Gly Leu Thr Asp Asp Gly Glu Ala Arg Val Gln Ala Leu Gly Met 85 90 Thr Pro Gly Ile Ala Phe Ser *

103

<210> 1398 <211> 82 <212> PRT

100

<213> Homo sapiens

<210> 1399

<211> 68

<212> PRT

<213> Homo sapiens

<400> 1399

 Met Gly Ala Val Leu Leu Val Cys Leu Gln Thr Ser Ile Ala Ala Arg

 1
 5
 10
 15

 Asp Asp Leu Lys Asp Ala Val Asp Ser Gly Leu Leu Leu Ala Asn Ser
 20
 25
 30

 Leu Ser His Phe Val Pro Leu Val Val Arg Asn Tyr Leu Val His Cys
 45

 Asn Leu Leu Gln Thr Leu Lys Phe Leu Leu Gly Asn Cys Thr Ala Gly
 50
 55
 60

 Lys Ala Ser *

<210> 1400

<211> 54

65 67

<212> PRT

<213> Homo sapiens

<400> 1400

<210> 1401

<211> 232

<212> PRT

<213> Homo sapiens

<400> 1401

Met Leu Phe Ala Phe Ile Ser Leu Leu Val Met Leu Pro Thr Trp Trp 1 5 10 15

Ile Val Ser Ser Trp Leu Val Trp Gly Val Ile Leu Phe Val Tyr Leu

20 25 Val Ile Arg Ala Leu Arg Leu Trp Arg Thr Ala Lys Leu Gln Val Thr 40 Leu Lys Lys Tyr Ser Val His Leu Glu Asp Met Ala Thr Asn Ser Arg 55 Ala Phe Thr Asn Leu Val Arg Lys Ala Leu Arg Leu Ile Gln Glu Thr 75 Glu Val Ile Ser Arg Gly Phe Thr Leu Leu Leu Asp Arg Val Ser Ala 90 Ala Cys Pro Phe Asn Lys Ala Gly Gln His Pro Ser Gln His Leu Ile 105 Gly Leu Arg Lys Ala Val Tyr Arg Thr Leu Arg Ala Ser Phe Gln Ala 120 Ala Arg Leu Ala Thr Leu Tyr Met Leu Lys Asn Tyr Pro Leu Asn Ser 135 140 Glu Ser Asp Asn Val Thr Asn Tyr Ile Cys Val Val Pro Phe Lys Glu 150 155 Leu Gly Leu Gly Leu Ser Glu Glu Gln Ile Ser Glu Glu Glu Ala His 170 Lys Leu Tyr Arg Trp Leu Gln Pro Ala Cys Ile Glu Gly Phe Val Pro 185 Thr Leu Gly Gly Thr Glu Phe Arg Val Leu Gln Thr Val Ser Pro Ile 200 Thr Phe Tyr Ser Gln Phe Thr Ser Trp Ala Leu Thr Tyr Ser Ser Thr 215 Ser Ala Ser Ser Tyr Leu Ile * 230 231

<210> 1402

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1402

 Met
 Ala
 Pro
 Ala
 Pro
 Trp
 Leu
 Thr
 Pro
 Val
 Ile
 Pro
 Ala
 Leu

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 15
 15

 Trp
 Glu
 Ala
 Glu
 Asp
 Gly
 Ser
 Arg
 Gly
 Gln
 Glu
 Phe
 Lys
 Thr
 Ser

 Leu
 Ala
 Ser
 Met
 Val
 Lys
 Pro
 Arg
 Leu
 Tyr
 Tyr
 Lys
 Tyr
 Lys
 Asn
 *

 35
 47

<210> 1403

<211> 53

<212> PRT

<213> Homo sapiens

<400> 1403

Tyr Cys Pro His * 50 52

<210> 1404 <211> 90 <212> PRT <213> Homo sapiens

<210> 1405 <211> 477 <212> PRT <213> Homo sapiens

<400> 1405 Met Ala Gly Arg Gly Gly Ser Ala Leu Leu Ala Leu Cys Gly Ala Leu Ala Ala Cys Gly Trp Leu Leu Gly Ala Glu Ala Gln Glu Pro Gly Ala 25 Pro Ala Ala Gly Met Arg Arg Arg Arg Leu Gln Glu Asp Gly Ile Ser Phe Glu Tyr His Arg Tyr Pro Glu Leu Arg Glu Ala Leu Val 55 Ser Val Trp Leu Gln Cys Thr Ala Ile Ser Arg Ile Tyr Thr Val Gly 75 Arg Ser Phe Glu Gly Arg Glu Leu Leu Val Ile Glu Leu Ser Asp Asn 85 90 Pro Gly Val His Glu Pro Gly Glu Pro Glu Phe Lys Tyr Ile Gly Asn 105 Met His Gly Asn Glu Ala Val Gly Arg Glu Leu Leu Ile Phe Leu Ala 125 120 Gln Tyr Leu Cys Asn Glu Tyr Gln Lys Gly Asn Glu Thr Ile Val Asn 135 140 Leu Ile His Ser Thr Arg Ile His Ile Met Pro Ser Leu Asn Pro Asp 150 155 Gly Phe Glu Lys Ala Ala Ser Gln Pro Gly Glu Leu Lys Asp Trp Phe 170 Val Gly Arg Ser Asn Ala Gln Gly Ile Asp Leu Asn Arg Asn Phe Pro 185 Asp Leu Asp Arg Ile Val Tyr Val Asn Glu Lys Glu Gly Gly Pro Asn

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200
Asn His Leu Leu Lys Asn Met Lys Lys Ile Val Asp Gln Asn Thr Lys
  210 215
                                    220
Leu Ala Pro Glu Thr Lys Ala Val Ile His Trp Ile Met Asp Ile Pro
225 230
                                235
Phe Val Leu Ser Ala Asn Leu His Gly Gly Asp Leu Val Ala Asn Tyr
            245
                             250
Pro Tyr Asp Glu Thr Arg Ser Gly Ser Ala His Glu Tyr Ser Ser Ser
                          265 270
Pro Asp Asp Ala Ile Phe Gln Ser Leu Ala Arg Ala Tyr Ser Ser Phe
     275
         280
Asn Pro Ala Met Ser Asp Pro Asn Arg Pro Pro Cys Arg Lys Asn Asp
                  295
Asp Asp Ser Ser Phe Val Asp Gly Thr Thr Asn Gly Gly Ala Trp Tyr
               310
                                315
Ser Val Pro Gly Gly Met Gln Asp Phe Asn Tyr Leu Ser Ser Asn Cys
                   330 335
            325
Phe Glu Ile Thr Val Glu Leu Ser Cys Glu Lys Phe Pro Pro Glu Glu
      340 345
Thr Leu Lys Thr Tyr Trp Glu Asp Asn Lys Asn Ser Leu Ile Ser Tyr
              360
                          365
Leu Glu Gln Ile His Arg Gly Val Lys Gly Phe Val Arg Asp Leu Gln
   370 375
                          380
Gly Asn Pro Ile Ala Asn Ala Thr Ile Ser Val Glu Gly Ile Asp His
385 390 395 400
Asp Val Thr Ser Ala Lys Asp Gly Asp Tyr Trp Arg Leu Leu Ile Pro
                            410
Gly Asn Tyr Lys Leu Thr Ala Ser Ala Pro Gly Tyr Leu Ala Ile Thr
                         425
Lys Lys Val Ala Val Pro Tyr Ser Pro Ala Ala Gly Val Asp Phe Glu
Leu Glu Ser Phe Ser Glu Arg Lys Glu Glu Glu Lys Glu Glu Leu Met
                  455
Glu Trp Trp Lys Met Met Ser Glu Thr Leu Asn Phe *
                470
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<210> 1406

<211> 55

<212> PRT

<213> Homo sapiens

<400> 1406

<210> 1407

<211> 66

<212> PRT

<213> Homo sapiens

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<210> 1409 <211> 72 <212> PRT <213> Homo sapiens

70 71

<210> 1410 <211> 53 <212> PRT <213> Homo sapiens

<210> 1411 <211> 82 <212> PRT <213> Homo sapiens

<400> 1411 Met Ala Ser Gln Ser Met Cys Phe Leu Trp Leu Ala Pro Val Thr Trp 1 5 10 Cys Val Met Phe Ser Ser Arg Thr Cys Tyr Ser Pro Cys Gly Asn Phe 20 25 Ser Thr Ala Pro Gly Arg Val Ile Phe His Ser Trp Asp Arg Ala Gln 40 Phe Val Tyr Ser Phe Leu Ser Arg Trp Arg Leu Gly Leu Phe Pro Pro 55 60 Leu Ala Ser Val Asn Gly Asp Ala Val Ile Met Gly Val Pro Val Phe 70 Val * 81

<210> 1412 <211> 72 <212> PRT <213> Homo sapiens

<210> 1413 <211> 59 <212> PRT

<213> Homo sapiens

<400> 1413

Met Met Thr Ile Lys Glu Phe Thr Leu Leu Leu Val Ser Leu Gln Phe 10 Ser Thr Phe Pro Ser Lys Lys Phe Leu Leu Glu Thr His Phe Leu Lys 25 Asn Ser Glu Asn Trp Leu Gly Val Val Ala His Ala Cys Ser Leu Ser 40 Thr Leu Gly Trp Pro Arg Arg Arg Thr Ala * 55

<210> 1414

<211> 78

<212> PRT

<213> Homo sapiens

<400> 1414

Met Leu Arg Leu Asp Ile Ile Asn Ser Leu Val Thr Thr Val Phe Met 10 Leu Ile Val Ser Val Leu Ala Leu Ile Pro Glu Thr Thr Thr Leu Thr 25 Val Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 40 Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Asn Pro Ser Gly Pro 55 Tyr Gln Lys Lys Pro Val His Glu Lys Lys Glu Val Leu * 70

<210> 1415 <211> 171 <212> PRT

<213> Homo sapiens

<400> 1415

Met His Met Met Lys Leu Ser Ile Lys Val Leu Leu Gln Ser Ala Leu 10 Ser Leu Gly Arg Ser Leu Asp Ala Asp His Ala Pro Leu Gln Gln Phe 25 Phe Val Val Met Glu His Cys Leu Lys His Gly Leu Lys Val Lys 40 Ser Phe Ile Gly Gln Asn Lys Ser Phe Phe Gly Pro Leu Glu Leu Val 55 Glu Lys Leu Cys Pro Glu Ala Ser Asp Ile Ala Thr Ser Val Arg Asn 70 Leu Pro Glu Leu Lys Thr Ala Val Gly Arg Gly Arg Ala Trp Leu Tyr 90 Leu Ala Leu Met Gln Lys Lys Leu Ala Asp Tyr Leu Lys Val Leu Ile 105 Asp Asn Lys His Leu Leu Ser Glu Phe Tyr Glu Pro Glu Ala Leu Met 120 Met Glu Glu Gly Met Val Ile Val Gly Leu Leu Val Gly Leu Asn

130 135 140

Val Leu Asp Ala Asn Leu Trp Leu Glu Arg Arg Arg Leu Gly Phe Ser

145 150 155 160

Gly Trp Ser Asn Arg Phe Phe Pro Leu Pro *

165 170

<210> 1416 <211> 77 <212> PRT <213> Homo sapiens

<210> 1417 <211> 249 <212> PRT <213> Homo sapiens

<400> 1417 Met Glu Lys Ile Pro Glu Ile Gly Lys Phe Gly Glu Lys Ala Pro Pro Ala Pro Ser His Val Trp Arg Pro Ala Ala Leu Phe Leu Thr Leu Leu Cys Leu Leu Leu Ile Gly Leu Gly Val Leu Ala Ser Met Phe His Val Thr Leu Lys Ile Glu Met Lys Lys Met Asn Lys Leu Gln Asn Ile Ser Glu Glu Leu Gln Arg Asn Ile Ser Leu Gln Leu Met Ser Asn Met 75 Asn Ile Ser Asn Lys Ile Arg Asn Leu Ser Thr Thr Leu Gln Thr Ile 90 Ala Thr Lys Leu Cys Arg Glu Leu Tyr Ser Lys Glu Gln Glu His Lys 105 Cys Lys Pro Cys Pro Arg Arg Trp Ile Trp His Lys Asp Ser Cys Tyr 120 Phe Leu Ser Asp Asp Val Gln Thr Trp Gln Glu Ser Lys Met Ala Cys 135 140 Ala Ala Gln Asn Ala Ser Leu Leu Lys Ile Asn Asn Lys Asn Ala Leu 155 150 Glu Phe Ile Lys Ser Gln Ser Arg Ser Tyr Asp Tyr Trp Leu Gly Leu 165 170 175 Ser Pro Glu Glu Asp Ser Thr Arg Gly Met Arg Val Asp Asn Ile Ile 180 185

Asn Ser Ser Ala Trp Val Ile Arg Asn Ala Pro Asp Leu Asn Asn Met
195

Tyr Cys Gly Tyr Ile Asn Arg Leu Tyr Val Gln Tyr Tyr His Cys Thr
210

Tyr Lys Gln Arg Met Ile Cys Glu Lys Met Ala Asn Pro Val Gln Leu
225

Gly Ser Thr Tyr Phe Arg Glu Ala *
245

Asn Asn Pro Val Gln Leu
248

<210> 1418 <211> 65 <212> PRT <213> Homo sapiens

<210> 1419 <211> 468 <212> PRT <213> Homo sapiens

<400> 1419 Met Leu Leu Leu Leu Leu Pro Leu Leu Trp Gly Arg Glu Arg Val Glu Gly Gln Lys Ser Asn Arg Lys Asp Tyr Ser Leu Thr Met Gln Ser 20 25 Ser Val Thr Val Gln Glu Gly Met Cys Val His Val Arg Cys Ser Phe 40 Ser Tyr Pro Val Asp Ser Gln Thr Asp Ser Asp Pro Val His Gly Tyr 55 Trp Phe Arg Ala Gly Asn Asp Ile Ser Trp Lys Ala Pro Val Ala Thr 75 70 Asn Asn Pro Ala Trp Ala Val Gln Glu Glu Thr Arg Asp Arg Phe His 90 85 Leu Leu Gly Asp Pro Gln Thr Lys Asn Cys Thr Leu Ser Ile Arg Asp 105 Ala Arg Met Ser Asp Ala Gly Arg Tyr Phe Phe Arg Met Glu Lys Gly 125 120 115 Asn Ile Lys Trp Asn Tyr Lys Tyr Asp Gln Leu Ser Val Asn Val Thr 140 135 Ala Leu Thr His Arg Pro Asn Ile Leu Ile Pro Gly Thr Leu Glu Ser 150 155 Gly Cys Phe Gln Asn Leu Thr Cys Ser Val Pro Trp Ala Cys Glu Gln

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170
               165
Gly Thr Pro Pro Met Ile Ser Trp Met Gly Thr Ser Val Ser Pro Leu
                              185
His Pro Ser Thr Thr Arg Ser Ser Val Leu Thr Leu Ile Pro Gln Pro
                           200
Gln His His Gly Thr Ser Leu Thr Cys Gln Val Thr Leu Pro Gly Ala
                       215
Gly Val Thr Thr Asn Arg Thr Ile Gln Leu Asn Val Ser Tyr Pro Pro
                   230
                                       235
Gln Asn Leu Thr Val Thr Val Phe Gln Gly Glu Gly Thr Ala Ser Thr
               245
                                  250
Ala Leu Gly Asn Ser Ser Ser Leu Ser Val Leu Glu Gly Gln Ser Leu
                 265
Arg Leu Val Cys Ala Val Asp Ser Asn Pro Pro Ala Arg Leu Ser Trp
                          280
                                             285
Thr Trp Arg Ser Leu Thr Leu Tyr Pro Ser Gln Pro Ser Asn Pro Leu
                                   300
                      295
Val Leu Glu Leu Gln Val His Leu Gly Asp Glu Gly Glu Phe Thr Cys
                  310
                                      315
Arg Ala Gln Asn Ser Leu Gly Ser Gln His Val Ser Leu Asn Leu Ser
              325
                                  330
Leu Gln Gln Glu Tyr Thr Gly Lys Met Arg Pro Val Ser Gly Val Leu
                              345
Leu Gly Ala Val Gly Gly Ala Gly Ala Thr Ala Leu Val Phe Leu Ser
                           360
Phe Cys Val Ile Phe Ile Val Val Arg Ser Cys Arg Lys Lys Ser Ala
                       375
Arg Pro Ala Ala Asp Val Gly Asp Ile Gly Met Lys Asp Ala Asn Thr
                  390
                                       395
Ile Arg Gly Ser Ala Ser Gln Gly Asn Leu Thr Glu Ser Trp Ala Asp
                                  410
Asp Asn Pro Arg His His Gly Leu Ala Ala His Ser Ser Gly Glu Glu
           420
                              425
                                                  430
Arg Glu Ile Gln Tyr Ala Pro Leu Ser Phe His Lys Gly Glu Pro Gln
                          440
                                              445
Asp Leu Ser Gly Gln Glu Ala Thr Asn Asn Glu Tyr Ser Glu Ile Lys
                      455
Ile Pro Lys *
     467
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<210> 1420 <211> 150 <212> PRT

<213> Homo sapiens

<210> 1421 <211> 89 <212> PRT <213> Homo sapiens

<400> 1421 Met Tyr Val Phe Leu Cys Pro Ala Cys Gly Arg Leu Met Gly Ser 10 Thr Tyr Met Arg Leu Leu Pro Gln Ser Glu Pro Ala Leu His Asn Arg 20 25 Ile Leu Arg Gln Thr Glu Pro Leu Leu Tyr Phe Lys Arg Gly Lys Gln 40 Gln Gly Leu Phe Tyr Ala Ser Phe Pro Ala Val His Arg Met Asp Ser 60 55 Leu Leu Arg Arg Thr Val Val Ile Leu Tyr Lys Arg Thr Asn Thr Val 70 75 Gly Val Ser Leu Phe Gln Asn Ala * 85

<210> 1422 <211> 83 <212> PRT <213> Homo sapiens

<400> 1422 Met Met Thr Trp Ala Ser Leu Ala Leu Gly Leu Thr Arg Ala Leu Gly 10 Gly Met Gly Ser Phe Leu Leu Arg Ile Leu Gly Trp Ser Trp Ala Met 20 25 Gly Ser Arg Ser Arg Ala Arg Trp Pro Arg Gly Arg Leu Gly Phe Thr 40 Ser Met Leu Ser Cys Met Arg Gln Cys Ser Val Cys Arg Met Ile Met 60 55 Ser Leu Val Glu Val Leu Val Ala Thr Ser Gln Val Val Lys Leu Trp 65 70 75 Ser Arg * 82

<210> 1423 <211> 54

<212> PRT <213> Homo sapiens

<210> 1424 <211> 73 <212> PRT <213> Homo sapiens

<210> 1425 <211> 245 <212> PRT <213> Homo sapiens

<400> 1425 Met Ala Cys Tyr Leu Leu Val Ala Asn Ile Leu Leu Val Asn Leu Leu 10 Ile Ala Val Phe Asn Asn Thr Phe Phe Glu Val Lys Ser Ile Ser Asn 25 Gln Val Trp Lys Phe Gln Arg Tyr Gln Leu Ile Met Thr Phe His Glu 40 Arg Pro Val Leu Pro Pro Pro Leu Ile Ile Phe Ser His Met Thr Met 55 Ile Phe Gln His Leu Cys Cys Arg Trp Arg Lys His Glu Ser Asp Pro 70 Asp Glu Arg Asp Tyr Gly Leu Lys Leu Phe Ile Thr Asp Asp Glu Leu 85 90 Lys Lys Val His Asp Phe Glu Glu Gln Cys Ile Glu Glu Tyr Phe Arg 105 Glu Lys Asp Asp Arg Phe Asn Ser Ser Asn Asp Glu Arg Ile Arg Val 120 125

Thr Ser Glu Arg Val Glu Asn Met Ser Met Arg Leu Glu Glu Val Asn 135 140 Glu Arg Glu His Ser Met Lys Ala Ser Leu Gln Thr Val Asp Ile Arg 150 155 Leu Ala Gln Leu Glu Asp Leu Ile Gly Arg Met Ala Thr Ala Leu Glu 165 170 Arg Leu Thr Gly Leu Glu Arg Ala Glu Ser Asn Lys Ile Arg Ser Arg 180 185 Thr Ser Ser Asp Cys Thr Asp Ala Arg Leu His Trp Pro Val Arg Ala 200 Ala Leu Thr Ser Gln Glu Arg Glu His Leu Ser Ala Pro Lys Arg Gly 215 220 Leu Glu Pro Trp Gln Asn Ile Leu Phe Ile Gln Tyr Lys Pro Ala Ala 230 235 Ser Ser Ser Thr 244

<210> 1426

<211> 520

<212> PRT

<213> Homo sapiens

<221> misc feature

<222> (1)...(520)

<223> Xaa = any amino acid or nothing

<400> 1426

Met Asp Ile Leu Leu Leu Leu Phe Phe Met Ile Ile Phe Ala Ile 5 10 Leu Gly Phe Tyr Leu Phe Ser Pro Asn Pro Ser Asp Pro Tyr Phe Ser 20 25 3.0 Thr Leu Glu Asn Ser Ile Val Ser Leu Phe Val Leu Leu Thr Thr Ala 4 O 45 Asn Phe Pro Asp Val Met Met Pro Ser Tyr Ser Arg Asn Pro Trp Ser 55 60 Cys Val Phe Phe Ile Val Tyr Leu Ser Ile Glu Leu Tyr Phe Ile Met 70 75 Asn Leu Leu Ala Val Val Phe Asp Thr Phe Asn Asp Ile Glu Lys 90 Arg Lys Phe Lys Ser Leu Leu His Lys Arg Thr Ala Ile Gln His 105 110 Ala Tyr Arg Leu Leu Ile Ser Glm Arg Arg Pro Ala Gly Ile Ser Tyr 120 125 Arg Gln Phe Glu Gly Leu Met Arg Phe Tyr Lys Pro Arg Met Ser Ala 135 140 Arg Glu Arg Tyr Leu Thr Phe Lys Ala Leu Asn Gln Asn Asn Thr Pro 150 155 Leu Leu Ser Leu Lys Asp Phe Tyr Asp Ile Tyr Glu Val Ala Ala Leu 165 170 Lys Trp Lys Ala Thr Lys Asn Arg Glu His Trp Val Asp Glu Leu Pro 180 185 Arg Thr Ala Leu Leu Ile Phe Lys Gly Ile Asn Ile Leu Val Lys Ala 200 Lys Ala Phe Gln Tyr Phe Met Tyr Leu Val Val Ala Val Asn Gly Val 215 220 Trp Ile Leu Val Glu Thr Phe Met Leu Lys Gly Gly Asn Phe Phe Ser

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225
                 230
                                  235
Lys His Val Pro Trp Ser Tyr Leu Val Phe Leu Thr Ile Tyr Gly Val .
             245
                              250
Glu Leu Phe Leu Lys Val Ala Gly Leu Gly Pro Val Glu Tyr Leu Ser
                           265
Ser Gly Trp Asn Leu Phe Asp Phe Ser Val Thr Val Phe Ala Phe Leu
             280
                                        285
Gly Leu Leu Ala Leu Ala Leu Asn Met Glu Pro Phe Tyr Phe Ile Val
        295
                                      300
Val Leu Arg Pro Leu Gln Leu Leu Arg Leu Phe Lys Leu Lys Glu Arg
                 310
                                  315
Tyr Arg Asn Val Leu Asp Thr Met Phe Glu Leu Leu Pro Arg Met Ala
             325
                               330
Ser Leu Gly Leu Thr Leu Leu Ile Phe Tyr Tyr Ser Phe Ala Ile Val
                           345
Gly Met Glu Phe Phe Cys Gly Ile Val Phe Pro Asn Cys Cys Asn Thr
                        360
                                         365
Ser Thr Val Ala Asp Ala Tyr Arg Trp Arg Asn His Thr Val Gly Asn
                    375
                               380
Arg Thr Val Val Glu Glu Gly Tyr Tyr Tyr Leu Asn Asn Phe Asp Asn
                                  395
                 390
Ile Leu Asn Ser Phe Val Thr Leu Phe Glu Leu Thr Val Val Asn Asn
             405
                              410
Trp Tyr Ile Ile Met Glu Gly Val Thr Ser Gln Thr Ser His Trp Ser
                 425
Arg Leu Tyr Phe Met Thr Phe Tyr Ile Ala Thr Met Val Val Met Thr
435 440
                                        445
Ile Ile Val Ala Phe Ile Leu Glu Ala Phe Val Phe Arg Met Asn Tyr
              455
                                     460
Ser Arg Lys Asn Gln Asp Ser Glu Val Asp Gly Gly Ile Thr Leu Glu
          470
                                 475 480
Lys Glu Ile Ser Lys Glu Glu Leu Val Ala Val Leu Glu Leu Tyr Arg
           485 490
Glu Ala Arg Xaa Ala Ser Ser Asp Val Thr Arg Leu Leu Glu Thr Leu
                          505
Ser Gln Met Glu Arg Tyr Gln Gln
      515
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<210> 1427 <211> 106 <212> PRT

<213> Homo sapiens

<400> 1427

 Met
 Ser
 Pro
 Gln
 His
 Leu
 Leu
 Leu
 Thr
 Leu
 Pro
 Leu
 Pro
 Leu
 Pro
 Leu
 Arg
 Ser

 Pro
 Ile
 Leu
 Phe
 Ser
 His
 Thr
 Ala
 Gln
 Leu
 Leu
 Leu
 Thr
 Arg
 Ile

 Ala
 Phe
 Arg
 Ala
 Cys
 Glu
 Leu
 Phe
 Phe
 Phe
 Val
 Met
 Val
 Ser
 Leu
 Cys

 Ala
 Phe
 Arg
 Ala
 His
 Ser
 Phe
 Ile
 Ala
 Thr
 Ile
 Thr
 Tyr
 Glu
 Arg
 Arg

 Cys
 Pro
 Gly
 Ile
 Ser
 Ser
 Phe
 Ile
 Ala
 Thr
 Ile
 Thr
 Tyr
 Glu
 Arg
 Arg

 Ala
 Phe
 Glu
 Glu
 Glu
 Fhe
 Ser
 Pro
 Pro
 Arg
 Glu

Thr Thr His Arg Leu Pro Ser Cys Phe * 100 105

<210> 1428 <211> 841 <212> PRT <213> Homo sapiens

<400> 1428 Met Ala Leu Ala Ser Ala Ala Pro Gly Ser Ile Phe Cys Lys Gln Leu 10 Leu Phe Ser Leu Leu Val Leu Thr Leu Leu Cys Asp Ala Cys Gln Lys 25 Val Tyr Leu Arg Val Pro Ser His Leu Gln Ala Glu Thr Leu Val Gly 40 Lys Val Asn Leu Glu Glu Cys Leu Lys Ser Ala Ser Leu Ile Arg Ser 55 60 Ser Asp Pro Ala Phe Arg Ile Leu Glu Asp Gly Ser Ile Tyr Thr Thr 75 His Asp Leu Ile Leu Ser Ser Glu Arg Lys Ser Phe Ser Ile Phe Leu 90 Ser Asp Gly Gln Arg Arg Glu Gln Glu Ile Lys Val Val Leu Ser 105 Ala Arg Glu Asn Lys Ser Pro Lys Lys Arg His Thr Lys Asp Thr Ala 120 125 Leu Lys Arg Ser Lys Arg Arg Trp Ala Pro Ile Pro Ala Ser Leu Met 135 140 Glu Asn Ser Leu Gly Pro Phe Pro Gln His Val Gln Gln Ile Gln Ser 150 155 Asp Ala Ala Gln Asn Tyr Thr Ile Phe Tyr Ser Ile Ser Gly Pro Gly 165 170 Val Asp Lys Glu Pro Phe Asn Leu Phe Tyr Ile Glu Lys Asp Thr Gly 185 Asp Ile Phe Cys Thr Arg Ser Ile Asp Arg Glu Lys Tyr Glu Gln Phe 200 205 Ala Leu Tyr Gly Tyr Ala Thr Thr Ala Asp Gly Tyr Ala Pro Glu Tyr 215 220 Pro Leu Pro Leu Ile Ile Lys Ile Glu Asp Asp Asn Asp Asn Ala Pro 230 235 Tyr Phe Glu His Arg Val Thr Ile Phe Thr Val Pro Glu Asn Cys Arg 245 250 Ser Gly Thr Ser Val Gly Lys Val Thr Ala Thr Asp Leu Asp Glu Pro 260 265 Asp Thr Leu His Thr Arg Leu Lys Tyr Lys Ile Leu Gln Gln Ile Pro 280 Asp His Pro Lys His Phe Ser Ile His Pro Asp Thr Gly Val Ile Thr 295 Thr Thr Pro Phe Leu Asp Arg Glu Lys Cys Asp Thr Tyr Gln Leu 315 Ile Met Glu Val Arg Asp Met Gly Gly Gln Pro Phe Gly Leu Phe Asn 325 330 Thr Gly Thr Ile Thr Ile Ser Leu Glu Asp Glu Asn Asp Asn Pro Pro 345 Ser Phe Thr Glu Thr Ser Tyr Val Thr Glu Val Glu Glu Asn Arg Ile 360 Asp Val Glu Ile Leu Arg Met Lys Val Gln Asp Gln Asp Leu Pro Asn

	370					375					380				
Thr 385	Pro	His	Ser	Lys	Ala 390		Tyr	Lys	Ile	Leu 395			Asn	Glu	Asn 400
Gly	Asn	Phe	Ile	Ile 405	Ser	Thr	Asp	Pro	Asn 410	Thr	Asn	Glu	Gly	Val 415	
			420				Tyr	425					430		
Gln	Val	Gly 435	Val	Ile	Asn	Glu	Ala 440	Gln	Phe	Ser	Lys	Ala 445	Ala	Ser	Ser
	450					455	Thr				460				_
465					470		His			475	-				480
	•			485			Gln		490					495	
			500				Glu	505					510		
		515					Ile 520					525	=		
	530					535	Glu				540				
545					550		Gly			555				_	560
				565			Asp		570					575	
			580				Cys	585					590		
		595					Gly 600					605			
	610					615	Ser	_		_	620		_	-	_
625					630		Arg			635					640
				645			Ile		650					655	
			660				Val	665					670		
		675		_	=		Thr 680	_	_			685			
	690					695	Ala				700				
705					710		Phe			715					720
				725			Ile		730					735	
			740				Glu	745					750	_	
•		755					Cys 760					765			
	770					775	Thr				780				
785					790		Asn			795		•			800
				805			Gln		810					815	
			820				Gln	Pro 825	Arg	Leu	Gly	Glu	Glu 830	Ser	Ile
Arg	Gly	His 835	Thr	Leu	Ile	Lys	Asn 840	*							

55

Ile Thr Asn Glu Leu Leu Lys Met Ser Leu Met Ser Thr Ala Thr Phe

Lys Asn Asp Ser Ile Ile Ser Asn Val Thr Val Thr Ser Val Thr Leu 100 105 110

Pro Asn Ala Val Ser Thr Leu Gln Ser Ser Lys Pro Lys Thr Glu Thr 115 120 125

Gln Ser Ser Ile Lys Thr Thr Glu Ile Pro Gly Ser Val Leu Gln Pro 130 135 140

Asp Ala Ser Pro Ser Lys Thr Gly Thr Leu Thr Ser Ile Pro Val Thr
145 150 155 160

Ile Pro Glu Asn Thr Ser Gln Ser Gln Val Ile Gly Thr Glu Gly Gly

165
170
175
175

Lys Asn Ala Ser Thr Ser Ala Thr Ser Arg Ser Tyr Ser Ser Ile Ile
180
185
190

Leu Pro Val Val Ile Ala Leu Ile Val Ile Thr Leu Ser Val Phe Val 195 200 205

Leu Val Gly Leu Tyr Arg Met Cys Trp Lys Ala Asp Pro Gly Thr Pro
210 225 220

Glu Asn Gly Asn Asp Gln Pro Gln Ser Asp Lys Glu Ser Val Lys Leu 225 230 235 240

Leu Thr Val Lys Thr Ile Ser His Glu Ser Gly Glu His Ser Ala Gln
245 250 255

Gly Lys Thr Lys Asn * 260 261

<210> 1429 <211> 262

<210> 1430

<211> 66

<212> PRT

<213> Homo sapiens

<400> 1430

Met Ser Tyr Thr Ala Phe Leu Ser Val Cys Cys Leu Pro Leu Leu Pro

1 5 10 15

Leu Cys Asp Phe Ala Leu Tyr Val Leu Leu Asp Lys Phe Lys Gly Gly

20 25 30

Phe Arg Gln Gln Asn Ser Pro Gln Ser Ile Tyr Gln His Asn Pro Tyr

35 40 45

Gln Asn Pro Asn Asn Val Leu Ile Phe Leu Gln Lys Trp Lys Asn Arg
50 55 60

Cys *
65

<210> 1431 <211> 437 <212> PRT <213> Homo sapiens

<400> 1431 Met Leu Lys Val Ser Ala Val Leu Cys Val Cys Ala Ala Ala Trp Cys 10 Ser Gln Ser Leu Ala Ala Ala Ala Ala Val Ala Ala Gly Gly Arg 25 Ser Asp Gly Gly Asn Phe Leu Asp Asp Lys Gln Trp Leu Thr Thr Ile Ser Gln Tyr Asp Lys Glu Val Gly Gln Trp Asn Lys Phe Arg Asp Glu Val Glu Asp Asp Tyr Phe Arg Thr Trp Ser Pro Gly Lys Pro Phe Asp 70 Gln Ala Leu Asp Pro Ala Lys Asp Pro Cys Leu Lys Met Lys Cys Ser 90 Arg His Lys Val Cys Ile Ala Gln Asp Ser Gln Thr Ala Val Cys Ile 105 Ser His Arg Arg Leu Thr His Arg Met Lys Glu Ala Gly Val Asp His 120 125 Arg Gln Trp Arg Gly Pro Ile Leu Ser Thr Cys Lys Gln Cys Pro Val 135 140 Val Tyr Pro Ser Pro Val Cys Gly Ser Asp Gly His Thr Tyr Ser Phe 150 155 Gln Cys Lys Leu Glu Tyr Gln Ala Cys Val Leu Gly Lys Gln Ile Ser 170 Val Lys Cys Glu Gly His Cys Pro Cys Pro Ser Asp Lys Pro Thr Ser 185 Thr Ser Arg Asn Val Lys Arg Ala Cys Ser Asp Leu Glu Phe Arg Glu 200 205 Val Ala Asn Arg Leu Arg Asp Trp Phe Lys Ala Leu His Glu Ser Gly 215 220 Ser Gln Asn Lys Lys Thr Lys Thr Leu Leu Arg Pro Glu Arg Ser Arg 230 235 Phe Asp Thr Ser Ile Leu Pro Ile Cys Lys Asp Ser Leu Gly Trp Met 245 250 Phe Asn Arg Leu Asp Thr Asn Tyr Asp Leu Leu Asp Gln Ser Glu 265 Leu Arg Ser Ile Tyr Leu Asp Lys Asn Glu Gln Cys Thr Lys Ala Phe 280 Phe Asn Ser Cys Asp Thr Tyr Lys Asp Ser Leu Ile Ser Asn Asn Glu 295 300 Trp Cys Tyr Cys Phe Gln Arg Gln Gln Asp Pro Pro Cys Gln Thr Glu 310 315 Leu Ser Asn Ile Gln Lys Arg Gln Gly Val Lys Lys Leu Leu Gly Gln 325 330 Tyr Ile Pro Leu Cys Asp Glu Asp Gly Tyr Tyr Lys Pro Thr Gln Cys

His Gly Ser Val Gly Gln Cys Trp Cys Val Asp Arg Tyr Gly Asn Glu 355 360 Val Met Gly Ser Arg Ile Asn Gly Val Ala Asp Cys Ala Ile Asp Phe 380 375 Glu Ile Ser Gly Asp Phe Ala Ser Gly Asp Phe His Glu Trp Thr Asp 395 390 Asp Glu Asp Asp Glu Asp Asp Ile Met Asn Asp Glu Asp Glu Ile Glu 410 Asp Asp Asp Glu Asp Glu Gly Asp Asp Asp Gly Gly Asp Asp His 425 420 Asp Val Tyr Ile * 435 436

<210> 1432

<211> 53

<212> PRT

<213> Homo sapiens

<400> 1432

Met Ser Tyr Val Glu Ile Leu Ile Pro Val Leu Leu Cys Leu His Ala 1 5 10 15

Phe Phe Pro Ser Ser Arg Arg His Val Ala Trp Phe Leu Ile Phe Ile 20 25 30

Cys Lys Phe Phe Lys Phe Cys Leu Ile Leu Lys Phe Ile Ile 11e Leu Ile 35

Leu Asn Tyr Leu * 50 52

<210> 1433

<211> 76

<212> PRT

<213> Homo sapiens

<400> 1433

 Met
 Glu
 Leu
 Lys
 Gly
 Phe
 Trp
 Leu
 Cys
 Leu
 Phe
 Leu
 Arg
 Phe
 Val
 Lys

 Trp
 Phe
 Val
 Asn
 Lys
 Gly
 Met
 Ile
 Leu
 Cys
 Thr
 Leu
 Phe
 Tyr
 Asn
 Leu

 Ile
 Tyr
 Ser
 Leu
 Tyr
 Asn
 Met
 Cys
 Trp
 Thr
 Val
 Leu
 Trp
 Ile
 Arg
 Lys

 Tyr
 Gln
 Thr
 Leu
 Leu
 Lys
 Glu
 Ser
 Phe
 Phe
 Ser
 Leu
 Asn
 Thr
 Phe
 Leu

 50
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<210> 1434

<211> 169

<212> PRT

<213> Homo sapiens

<400> 1434 Met Glu Ser Trp Trp Gly Leu Pro Cys Leu Ala Phe Leu Cys Phe Leu 10 Met His Ala Arg Gly Gln Arg Asp Phe Asp Leu Ala Asp Ala Leu Asp 25 Asp Pro Glu Pro Thr Lys Lys Pro Asn Ser Asp Ile Tyr Pro Lys Pro 40 Lys Pro Pro Tyr Tyr Pro Gln Pro Glu Asn Pro Asp Ser Gly Gly Asn 55 60 Ile Tyr Pro Arg Pro Lys Pro Arg Pro Gln Pro Gln Pro Gly Asn Ser 70 Gly Asn Ser Gly Gly Ser Tyr Phe Asn Asp Val Asp Arg Asp Asp Gly 90 Arg Tyr Pro Pro Arg Pro Arg Pro Arg Pro Pro Ala Gly Gly Gly Gly 105 Gly Gly Tyr Ser Ser Tyr Gly Asn Ser Asp Asn Thr His Gly Gly Asp 120 His His Ser Thr Tyr Gly Asn Pro Glu Gly Asn Met Val Ala Lys Ile 135 140 Val Ser Pro Ile Val Ser Val Val Val Thr Leu Leu Gly Ala Ala 150 155 Ala Gln Leu Phe Gln Thr Lys Gln * 165

<210> 1435 <211> 162 <212> PRT <213> Homo sapiens

<400> 1435

Met Arg Phe Val Thr Leu Ser Ser Ala Cys Leu Cys Pro Cys Pro Leu 1 5 10 Gly Pro Cys Trp Thr Arg His Pro Ser Tyr Gly Asn Leu His Glu Ala 25 Ser Thr Ser Leu Pro Pro Arg His Trp Thr Gly Ala Arg Lys Trp Asn 40 Glu Ser Ser His Cys Leu Lys Ser Trp Arg Pro Ser Ser Ala Ser Gly 55 60 Ser Pro Glu Asn Leu Gly Ser Asp Arg Arg Thr Glu Thr Glu Gly Arg 75 Glu Arg Asp Cys Asp Arg Glu Ala Glu Glu Gly Asp Arg Val Arg Glu 85 Glu Gln Asn Ser Leu Gln Trp Glu Gln Arg Gln Lys Cys Gly Gly Pro 105 Thr Gly Arg Gly Gly Arg Glu Gly Glu Gly Arg Arg Glu Gly Gln Leu 120 125 Pro Val Gln Val Ala Val Arg Ala Leu Gly Leu Gly Arg Gly Thr Leu 135 140 Leu Leu Ala Ser His Thr Gly Ser Ile Arg Gly Pro Arg Glu Gln 150 155 Val Ser 162

<210> 1436

<211> 77 <212> PRT <213> Homo sapiens

<210> 1437 <211> 85 <212> PRT <213> Homo sapiens

<210> 1438 <211> 76 <212> PRT <213> Homo sapiens

<210> 1439 <211> 425 <212> PRT <213> Homo sapiens

<400> 1439 Met Ser Leu Thr Ile Trp Thr Val Cys Gly Val Leu Ser Leu Phe Gly 10 Ala Leu Ser Tyr Ala Glu Leu Gly Thr Thr Ile Lys Lys Ser Gly Gly 25 His Tyr Thr Tyr Ile Leu Glu Val Phe Gly Pro Leu Pro Ala Phe Val 40 Arg Val Trp Val Glu Leu Leu Ile Ile Arg Pro Ala Ala Thr Ala Val 55 Ile Ser Leu Ala Phe Gly Arg Tyr Ile Leu Glu Pro Phe Phe Ile Gln 70 75 Cys Glu Ile Pro Glu Leu Ala Ile Lys Leu İle Thr Ala Val Gly Ile 90 85 Thr Val Val Met Val Leu Asn Ser Met Ser Val Ser Trp Ser Ala Arg 105 100 Ile Gln Ile Phe Leu Thr Phe Cys Lys Leu Thr Ala Ile Leu Ile Ile 125 120 Ile Val Pro Gly Val Met Gln Leu Ile Lys Gly Gln Thr Gln Asn Phe 135 140 Lys Asp Ala Phe Ser Gly Arg Asp Ser Ser Ile Thr Arg Leu Pro Leu 150 155 160 Ala Phe Tyr Tyr Gly Met Tyr Ala Tyr Ala Gly Trp Phe Tyr Leu Asn 170 Phe Val Thr Glu Glu Val Glu Asn Pro Glu Lys Thr Ile Pro Leu Ala 185 Ile Cys Ile Ser Met Ala Ile Val Thr Ile Gly Tyr Val Leu Thr Asn 200 Val Ala Tyr Phe Thr Thr Ile Asn Ala Glu Glu Leu Leu Ser Asn 215 Ala Val Ala Val Thr Phe Ser Glu Arg Leu Leu Gly Asn Phe Ser Leu 230 235 Ala Val Pro Ile Phe Val Ala Leu Ser Cys Phe Gly Ser Met Asn Gly 245 250 Gly Val Phe Ala Val Ser Arg Leu Phe Tyr Val Ala Ser Arg Glu Gly 265 His Leu Pro Glu Ile Leu Ser Met Ile His Val Arg Lys His Thr Pro 280 285 Leu Pro Ala Val Ile Val Leu His Pro Leu Thr Met Ile Met Leu Phe 295 300 Ser Gly Asp Leu Asp Ser Leu Leu Asn Phe Leu Ser Phe Ala Arg Trp 310 315 Leu Phe Ile Gly Leu Ala Val Ala Gly Leu Ile Tyr Leu Arg Tyr Lys 325 330 Cys Pro Asp Met His Arg Pro Phe Lys Val Pro Leu Phe Ile Pro Ala 345 Leu Phe Ser Phe Thr Cys Leu Phe Met Val Ala Leu Ser Leu Tyr Ser 360 Asp Pro Phe Ser Thr Gly Ile Gly Phe Val Ile Thr Leu Thr Gly Val 375 380 Pro Ala Tyr Tyr Leu Phe Ile Ile Trp Asp Lys Lys Pro Arg Trp Phe 390 395

PCT/US01/02687 WO 01/54477

Arg Ile Met Ser Glu Lys Ile Thr Arg Thr Leu Gln Ile Ile Leu Glu 410 405 Val Val Pro Glu Glu Asp Lys Leu * 420 424

<210> 1440 <211> 70 <212> PRT <213> Homo sapiens

<400> 1440 Met Ser Val Phe Trp Gly Phe Val Gly Phe Leu Val Pro Trp Phe Ile 10 Pro Lys Gly Pro Asn Arg Gly Val Ile Ile Thr Met Leu Val Thr Cys 25 Ser Val Cys Cys Tyr Leu Phe Trp Leu Ile Ala Ile Leu Ala Gln Leu 40 Asn Pro Leu Phe Gly Pro Gln Leu Lys Asn Glu Thr Ile Trp Tyr Leu 55 Lys Tyr His Trp Pro *

<210> 1441 <211> 1691 <212> PRT

<213> Homo sapiens

<400> 1441 Met Trp Ser Leu His Ile Val Leu Met Arg Cys Ser Phe Arg Leu Thr 10 Lys Ser Leu Ala Thr Gly Pro Trp Ser Leu Ile Leu Ile Leu Phe Ser 25 Val Gln Tyr Val Tyr Gly Ser Gly Lys Lys Tyr Ile Gly Pro Cys Gly 40 Gly Arg Asp Cys Ser Val Cys His Cys Val Pro Glu Lys Gly Ser Arg 55 Gly Pro Pro Gly Pro Gly Pro Gln Gly Pro Ile Gly Pro Leu Gly 75 Ala Pro Gly Pro Ile Gly Leu Ser Gly Glu Lys Gly Met Arg Gly Asp 90 Arg Gly Pro Pro Gly Ala Ala Gly Asp Lys Gly Asp Lys Gly Pro Thr 105 Gly Val Pro Gly Phe Pro Gly Leu Asp Gly Ile Pro Gly His Pro Gly 120 Pro Pro Gly Pro Arg Gly Lys Pro Gly Met Ser Gly His Asn Gly Ser 135 Arg Gly Asp Pro Gly Phe Pro Gly Gly Arg Gly Ala Leu Gly Pro Gly 150 155 Gly Pro Leu Gly His Pro Gly Glu Lys Gly Glu Lys Gly Asn Ser Val 165 170 Phe Ile Leu Gly Ala Val Lys Gly Ile Gln Gly Asp Arg Gly Asp Pro 185 180 Gly Leu Pro Gly Leu Pro Gly Ser Trp Gly Ala Gly Gly Pro Ala Gly

		195					200					205			
Pro	Thr 210	Gly	Tyr	Pro	Gly	Glu 215	Pro	Gly	Leu	Val	Gly 220	Pro	Pro	Gly	Gln
Pro 225	Gly	Arg	Pro	Gly	Leu 230	Lys	Gly	Asn	Pro	Gly 235	Val	Gly	Val	Lys	Gly 240
Gln	Met	Gly	Asp	Pro 245	Gly	Glu	Val	Gly	Gln 250	Gln	Gly	Ser	Pro	Gly 255	Pro
Thr	Leu	Leu	Val 260	Glu	Pro	Pro	Asp	Phe 265	Cys	Leu	Tyr	Lys	Gly 270	Glu	Lys
Gly	Ile	Lys 275	Gly	Ile	Pro	Gly	Met 280	Val	Gly	Leu	Pro	Gly 285	Pro	Pro	Gly
Arg	Lys 290	Gly	Glu	Ser	Gly	Ile 295	Gly	Ala	Lys	Gly	Glu 300	Lys	Gly	Ile	Pro
Gly 305	Phe	Pro	Gly	Pro	Arg 310	Gly	Asp	Pro	Gly	Ser 315	Tyr	Gly	Ser	Pro	Gly 320
Phe	Pro	Gly	Leu	Lys 325	Gly	Glu	Leu	Gly	Leu 330	Val	Gly	Asp	Pro	Gly 335	Leu
Phe	Gly	Leu	Ile 340	Gly	Pro	Lys	Gly	Asp 345	Pro	Gly	Asn	Arg	Gly 350	His	Pro
_		355	Gly				360					365	-	-	
	370		Pro			375					380				
385			Gly		390					395					400
			Ile	405					410					415	
Gly	Leu	Pro	Gly 420	Glu	Ala	Gly	Ile	Pro 425	Gly	Arg	Pro	Asp	Ser 430	Ala	Pro
		435	Gly	_		_	440		_			445			_
	450		Leu			455				-	460				
465			Gln	_	470	_	-	_		475			_	_	480
			Gly	485	_	_			490		_			495	
			Gly 500			_		505	_			_	510		_
		515	Asp				520					525			
	530		Pro			535					540				
545			Gly		550	_		_	_	555	_	_	_		560
Val	Ser	Arg	Val	Lys 565	Gly	His	Lys	Gly	Glu 570	'Arg	Gly	Pro	Asp	Gly 575	Pro
			Pro 580	_			_	585					590		
Gly	Glu	Lys 595	Gly	Asp	Pro	Gly	Pro 600	Pro	Gly	Asp	His	Glu 605	Asp	Ala	Thr
Pro	Gly 610	Gly	Lys	Gly	Phe	Pro 615	Gly	Pro	Leu	Gly	Pro 620	Pro	Gly	Lys	Ala
Gly 625	Pro	Val	Gly	Pro	Pro 630	Gly	Leu	Gly	Phe	Pro 635	Gly	Pro	Pro	Gly	Glu 640
			Pro	645			-		650				_	655	
Gly	Leu	Lys	Gly 660	Gln	Lys	Gly	Asp	Thr 665	Ile	Ser	Cys	Asn	Val 670	Thr	Tyr

Pro Gly Arg His Gly Pro Pro Gly Phe Asp Gly Pro Pro Gly Pro Lys Gly Phe Pro Gly Pro Gln Gly Ala Pro Gly Leu Ser Gly Ser Asp Gly His Lys Gly Arg Pro Gly Thr Pro Gly Thr Ala Glu Ile Pro Gly Pro Pro Gly Phe Arg Gly Asp Met Gly Asp Pro Gly Phe Gly Glu Lys Gly Ser Ser Pro Val Gly Pro Pro Gly Pro Pro Gly Ser Pro Gly Val Asn Gly Gln Lys Gly Ile Pro Gly Asp Pro Ala Phe Gly His Leu Gly Pro Pro Gly Lys Arg Gly Leu Ser Gly Val Pro Gly Ile Lys Gly Pro Arg Gly Asp Pro Gly Cys Pro Gly Ala Glu Gly Pro Ala Gly Ile Pro Gly Phe Leu Gly Leu Lys Gly Pro Lys Gly Arg Glu Gly His Ala Gly Phe Pro Gly Val Pro Gly Pro Pro Gly His Ser Cys Glu Arg Gly Ala Pro Gly Ile Pro Gly Gln Pro Gly Leu Pro Gly Tyr Pro Gly Ser Pro Gly Ala Pro Gly Gly Lys Gly Gln Pro Gly Asp Val Gly Pro Pro Gly Pro Ala Gly Met Lys Gly Leu Pro Gly Leu Pro Gly Arg Pro Gly Ala His Gly Pro Pro Gly Leu Pro Gly Ile Pro Gly Pro Phe Gly Asp Asp Gly Leu Pro Gly Pro Pro Gly Pro Lys Gly Pro Arg Gly Leu Pro Gly Phe Pro Gly Phe Pro Gly Glu Arg Gly Lys Pro Gly Ala Glu Gly Cys Pro Gly Ala Lys Gly Glu Pro Gly Glu Lys Gly Met Ser Gly Leu Pro Gly Asp Arg Gly Leu Arg Gly Ala Lys Gly Ala Ile Gly Pro Pro Gly Asp Glu Gly Glu Met Ala Ile Ile Ser Gln Lys Gly Thr Pro Gly Glu Pro Gly Pro Pro Gly Asp Asp Gly Phe Pro Gly Glu Arg Gly Asp Lys Gly Thr Pro Gly Met Gln Gly Arg Arg Gly Glu Leu Gly Arg Tyr Gly Pro Pro Gly Phe His Arg Gly Glu Pro Gly Glu Lys Gly Gln Pro Gly Pro Pro Gly Pro Pro Gly Pro Pro Gly Ser Thr Gly Leu Arg Gly Phe Ile Gly Phe Pro Gly Leu Pro Gly Asp Gln Gly Glu Pro Gly Ser Pro Gly Pro Pro Gly Phe Ser Gly Ile Asp Gly Ala Arg Gly Pro Lys Gly Asn Lys Gly Asp Pro Ala Ser His Phe Gly Pro Pro Gly Pro Lys Gly Glu Pro Gly Ser Pro Gly Cys Pro Gly His Phe Gly Ala Ser Gly Glu Gln Gly Leu Pro Gly Ile Gln Gly Pro Arg Gly Ser Pro Gly Arg Pro Gly Pro Pro Gly Ser Ser Gly Pro Pro Gly Cys Pro Gly Asp His Gly Met Pro Gly Leu Arg Gly Gln Pro Gly Glu Met Gly Asp Pro Gly Pro

1145 1150 Arg Gly Leu Gln Gly Asp Pro Gly Ile Pro Gly Pro Pro Gly Ile Lys 1160 1165 Gly Pro Ser Gly Ser Pro Gly Leu Asn Gly Leu His Gly Leu Lys Gly 1175 1180 Gln Lys Gly Thr Lys Gly Ala Ser Gly Leu His Asp Val Gly Pro Pro 1185 1190 1195 Gly Pro Val Gly Ile Pro Gly Leu Lys Gly Glu Arg Gly Asp Pro Gly 1205 1210 1215 Ser Pro Gly Ile Ser Pro Pro Gly Pro Arg Gly Lys Lys Gly Pro Pro 1220 1225 1230 Gly Pro Pro Gly Ser Ser Gly Pro Pro Gly Pro Ala Gly Ala Thr Gly 1235 1240 1245 Arg Ala Pro Lys Asp Ile Pro Asp Pro Gly Pro Pro Gly Asp Gln Gly 1250 1255 1260 Pro Pro Gly Pro Asp Gly Pro Arg Gly Ala Pro Gly Pro Pro Gly Leu 1265 1270 1275 Pro Gly Ser Val Asp Leu Leu Arg Gly Glu Pro Gly Asp Cys Gly Leu 1285 1290 1295 Pro Gly Pro Pro Gly Pro Pro Gly Pro Pro Gly Pro Pro Gly Tyr Lys 1300 1305 1310 Gly Phe Pro Gly Cys Asp Gly Lys Asp Gly Gln Lys Gly Pro Val Gly 1315 1320 1325 Phe Pro Gly Pro Gln Gly Pro His Gly Phe Pro Gly Pro Pro Gly Glu 1330 1335 1340 Lys Gly Leu Pro Gly Pro Pro Gly Arg Lys Gly Pro Thr Gly Leu Pro 1350 1355 1360 Gly Pro Arg Gly Glu Pro Gly Pro Pro Ala Asp Val Asp Asp Cys Pro 1365 1370 1375 Arg Ile Pro Gly Leu Pro Gly Ala Pro Gly Met Arg Gly Pro Glu Gly 1380 1385 1390 Ala Met Gly Leu Pro Gly Met Arg Gly Pro Ser Gly Pro Gly Cys Lys 1395 1400 1405 Gly Glu Pro Gly Leu Asp Gly Arg Arg Gly Val Asp Gly Val Pro Gly 1410 1415 1420 Ser Pro Gly Pro Pro Gly Arg Lys Gly Asp Thr Gly Glu Asp Gly Tyr 1425 1430 1435 1440 Pro Gly Gly Pro Gly Pro Gly Pro Ile Gly Asp Pro Gly Pro Lys 1445 1450 1455 Gly Phe Gly Pro Gly Tyr Leu Gly Gly Phe Leu Leu Val Leu His Ser 1460 1465 1470 Gln Thr Asp Gln Glu Pro Thr Cys Pro Leu Gly Met Pro Arg Leu Trp 1475 1480 1485 Thr Gly Tyr Ser Leu Leu Tyr Leu Glu Gly Gln Glu Lys Ala His Asn 1490 1495 1500 Gln Asp Leu Gly Leu Ala Gly Ser Cys Leu Pro Val Phe Ser Thr Leu 1505 1510 1515 1520 Pro Phe Ala Tyr Cys Asn Ile His Gln Val Cys His Tyr Ala Gln Arg 1525 1530 Asn Asp Arg Ser Tyr Trp Leu Ala Ser Ala Ala Pro Leu Pro Met Met 1540 1545 Pro Leu Ser Glu Glu Ala Ile Arg Pro Tyr Val Ser Arg Cys Ala Val 1560 1565 Cys Glu Ala Pro Ala Gln Ala Val Ala Val His Ser Gln Asp Gln Ser 1570 1575 1580 Ile Pro Pro Cys Pro Gln Thr Trp Arg Ser Leu Trp Ile Gly Tyr Ser 1585 1590 1595 Phe Leu Met His Thr Gly Ala Gly Asp Gln Gly Gly Gln Ala Leu 1605 1610

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Met Ser Pro Gly Ser Cys Leu Glu Asp Phe Arg Ala Ala Pro Phe Leu 1620 1625 1630 Glu Cys Gln Gly Arg Gln Gly Thr Cys His Phe Phe Ala Asn Lys Tyr 1635 1640 1645 Ser Phe Trp Leu Thr Thr Val Lys Ala Asp Phe Glu Phe Ser Ser Ala 1650 1655 1660 Pro Ala Pro Asp Thr Leu Lys Glu Ser Gln Ala Gln Arg Gln Lys Ile 1670 1675 Ser Arg Cys Gln Val Cys Val Lys Tyr Ser *

<210> 1442 <211> 153 <212> PRT <213> Homo sapiens

<400> 1442 Met Gly Val Met Ala Pro Arg Thr Leu Leu Leu Leu Leu Leu Gly Ala 5 10 Leu Ala Leu Thr Glu Thr Trp Ala Gly Glu Cys Gly Val Gly Arg Glu 25 Arg Ala Ser Ala Gly Arg Ser Glu Trp Pro Ala Arg Pro Gly Glu Pro Arg Arg Glu Glu Gly Arg Ala Gly Leu Ser Leu Ser Pro Pro Gly 55 Ser His Ser Leu Arg Tyr Phe Ser Thr Ala Val Ser Gln Pro Gly Arg 70 Gly Glu Pro Arg Phe Ile Ala Val Gly Tyr Val Asp Asp Thr Glu Phe 90 85 Val Arg Phe Asp Ser Asp Ser Val Ser Pro Arg Met Glu Arg Arg Ala 100 105 Pro Trp Val Glu Glu Glu Gly Leu Glu Tyr Trp Asp Gln Glu Thr Arg 120 Asn Ala Lys Gly His Ala Gln Ile Tyr Arg Val Asn Leu Arg Thr Leu 135 Leu Arg Tyr Tyr Asn Gln Ser Glu Ala 150 · 153

<210> 1443 <211> 58 <212> PRT <213> Homo sapiens

<400> 1443 Met Ser Leu Leu Cys Leu Lys Phe Phe Ser Gly Leu Trp Thr Ile Thr Phe Ser Lys Gly Ala Lys Ile Ile His Trp Gly Arg Ser Leu Phe Asn 25 Trp Ile Ser Met Cys Lys Arg Met Lys Leu Asp Pro Tyr Ser Tyr His 40 Thr Gln Lys Leu Thr Gln Asn Gly Ser * 50 55

<210> 1444 <211> 69 <212> PRT <213> Homo sapiens

<210> 1445 <211> 826 <212> PRT <213> Homo sapiens

<400> 1445 Met Gly Trp Leu Cys Ser Gly Leu Leu Phe Pro Val Ser Cys Leu Val 1 5 10 Leu Gln Val Ala Ser Ser Gly Asn Met Lys Val Leu Gln Glu Pro 20 25 Thr Cys Val Ser Asp Tyr Met Ser Ile Ser Thr Cys Glu Trp Lys Met 40 Asn Gly Pro Thr Asn Cys Ser Thr Glu Leu Arg Leu Leu Tyr Gln Leu 55 Val Phe Leu Leu Ser Glu Ala His Thr Cys Val Pro Glu Asn Asn Gly 70 75 Gly Ala Gly Cys Val Cys His Leu Leu Met Asp Asp Val Val Ser Ala 85 90 Asp Asn Tyr Thr Leu Asp Leu Trp Ala Gly Gln Gln Leu Leu Trp Lys 105 Gly Ser Phe Lys Pro Ser Glu His Val Lys Pro Arg Ala Pro Gly Asn 120 Leu Thr Val His Thr Asn Val Ser Asp Thr Leu Leu Leu Thr Trp Ser 135 140 Asn Pro Tyr Pro Pro Asp Asn Tyr Leu Tyr Asn His Leu Thr Tyr Ala 150 155 Val Asn Ile Trp Ser Glu Asn Asp Pro Ala Asp Phe Arg Ile Tyr Asn 165 170 Val Thr Tyr Leu Glu Pro Ser Leu Arg Ile Ala Ala Ser Thr Leu Lys 180 185 190 Ser Gly Ile Ser Tyr Arg Ala Arg Val Arg Ala Trp Ala Gln Cys Tyr 200 205 Asn Thr Thr Trp Ser Glu Trp Ser Pro Ser Thr Lys Trp His Asn Ser 215 220 Tyr Arg Glu Pro Phe Glu Gln His Leu Leu Gly Val Ser Val Ser 230 235

Cys	Ile	Val	Ile	Leu 245		Val	Cys	Leu	Leu 250		Tyr	Val	Ser	Ile 255	
Lys	Ile	Lys	Lys 260	Glu		Trp	Asp	Gln 265	Ile		Asn	Pro	Ala 270	Arg	
Arg	Leu	Val 275	Ala		Ile	Ile	Gln 280	Asp		Gln	Gly	Ser 285			Glu
Lys	Arg 290	_	Arg	Gly	Gln	Glu 295			Lys	Cys	Pro		Trp	Lys	Asn
Cys 305	Leu	Thr	Lys	Leu	Leu 310		Cys	Phe	Leu	Glu 315		Asn	Met	Lys	Arg
Asp	Glu	Asp	Pro	His		Ala	Ala	Lys	Glu 330		Pro	Phe	Gln	Gly 335	
Gly	Lys	Ser	Ala 340	Trp	Cys	Pro	Val	Glu 345		Ser	Lys	Thr	Val 350		Trp
Pro	Glu	Ser 355	Ile	Ser	Val	Val	Arg 360		Val	Glu	Leu	Phe 365		Ala	Pro
Val	Glu 370	Cys	Glu	Glu	Glu	Glu 375	Glu	Val	Glu	Glu	Glu 380	Lys	Gly	Ser	Phe
Cys 385		Ser	Pro	Glu	Ser 390	Ser	Arg	Asp	Asp	Phe 395	Gln	Glu	Gly	Arg	Glu 400
			Ala	405					410					415	
			Gly 420					425					430		
		435	Ser				440					445	_		
	450		Gly			455					460	_			
465			Glu		470					475					480
			Cys	485					490					495	
			Phe 500					505					510	_	
		515	Asp				520					525			
	530		Cys			535					540				
545			Glu		550					555					560
				565					570					575	
			His 580					585					590		
		595	Gly				600					605			
	610		Ser			615					620	_		_	
625			Glu		630					635					640
			Asp	645					650					655	
			Glu 660					665					670		
		675	Glu				680					685			
	690		Pro			695					700				
usb	Set	nen	Gly	ser	GTÀ	тте	vaı	ıyr	ser	АТА	ьeu	Inr	cys	Hls	ьeu

705 710 715 Cys Gly His Leu Lys Gln Cys His Gly Gln Glu Asp Gly Gly Gln Thr 725 730 Pro Val Met Ala Ser Pro Cys Cys Gly Cys Cys Gly Asp Arg Ala 745 Ser Pro Pro Thr Thr Pro Leu Arg Ala Pro Asp Pro Ser Pro Gly Gly 760 Val Pro Leu Glu Ala Ser Leu Cys Pro Ala Ser Leu Ala Pro Ser Gly 775 Ile Ser Glu Lys Ser Lys Ser Ser Ser Ser Phe His Pro Ala Pro Gly 790 795 Asn Ala Gln Ser Ser Gln Thr Pro Lys Ile Val Asn Phe Val Ser 810 805 Val Gly Pro Thr Tyr Met Arg Val Ser * 820

<210> 1446

<211> 367

<212> PRT

<213> Homo sapiens

<400> 1446 Met Ala Leu Arg Phe Leu Leu Gly Phe Leu Leu Ala Gly Val Asp Leu 1 5 Gly Val Tyr Leu Met Arg Leu Glu Leu Cys Asp Pro Thr Gln Arg Leu 25 Arg Val Ala Leu Ala Gly Glu Leu Val Gly Val Gly Gly His Phe Leu 40 45 Phe Leu Gly Leu Ala Leu Val Ser Lys Asp Trp Arg Phe Leu Gln Arg 55 Met Ile Thr Ala Pro Cys Ile Leu Phe Leu Phe Tyr Gly Trp Pro Gly 70 75 Leu Phe Leu Glu Ser Ala Arg Trp Leu Ile Val Lys Arg Gln Ile Glu 85 90 Glu Ala Gln Ser Val Leu Arg Ile Leu Ala Glu Arg Asn Arg Pro His 105 100 Gly Gln Met Leu Gly Glu Glu Ala Gln Glu Ala Leu Gln Asp Leu Glu 120 125 Asn Thr Cys Pro Leu Pro Ala Thr Ser Ser Phe Ser Phe Ala Ser Leu 135 140 Leu Asn Tyr Arg Asn Ile Trp Lys Asn Leu Leu Ile Leu Gly Phe Thr 150 155 Asn Phe Ile Ala His Ala Ile Arg His Cys Tyr Gln Pro Val Gly Gly 170 Gly Gly Ser Pro Ser Asp Phe Tyr Leu Cys Ser Leu Leu Ala Ser Gly 185 Thr Ala Ala Leu Ala Cys Val Phe Leu Gly Val Thr Val Asp Arg Phe 200 Gly Arg Arg Gly Ile Leu Leu Ser Met Thr Leu Thr Gly Ile Ala 215 . 220 Ser Leu Val Leu Leu Gly Leu Trp Asp Tyr Leu Asn Glu Ala Ala Ile 230 235 Thr Thr Phe Ser Val Leu Gly Leu Phe Ser Ser Gln Ala Ala Ala Ile 250 245 Leu Ser Thr Leu Leu Ala Ala Glu Val Ile Pro Thr Thr Val Arg Gly 260

Arg Gly Leu Gly Leu Ile Met Ala Leu Gly Ala Leu Gly Gly Leu Ser 275 280 · 285 Gly Pro Ala Gln Arg Leu His Met Gly His Gly Ala Phe Leu Gln His 295 300 Val Val Leu Ala Ala Cys Ala Leu Leu Cys Ile Leu Ser Ile Met Leu 310 315 Leu Pro Glu Thr Lys Arg Lys Leu Leu Pro Glu Val Leu Arg Asp Gly 325 330 Glu Leu Cys Arg Arg Pro Ser Leu Leu Arg Gln Pro Pro Pro Thr Arg 340 345 Cys Asp His Val Pro Leu Leu Ala Thr Pro Asn Pro Ala Leu * 355 360

<210> 1447

<211> 79

<212> PRT

<213> Homo sapiens

<400> 1447

 Met
 Ala
 Ile
 Ser
 Trp
 Leu
 Gly
 Thr
 Trp
 Leu
 Gln
 Ser
 His
 Arg
 His
 His
 15

 Trp
 Ser
 Glu
 Pro
 Gln
 Leu
 Cys
 Arg
 Leu
 Pro
 Ala
 Arg
 His
 Leu
 Ile
 30

 Asn
 Leu
 Asn
 Phe
 Met
 Val
 Ala
 Glu
 Gly
 Ile
 Gly
 Asp
 Arg
 Ala
 Trp
 His

 Ile
 Ser
 Ala
 Gln
 Leu
 Phe
 Met
 Thr
 Phe
 Ser
 Phe
 His
 Ala
 Val
 Ile

 50
 Ile
 Ser
 Ala
 Gly
 Gly
 Gly
 Ala
 Gly
 Fys
 Tyr
 Lys
 Asp
 Lys
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 65
 Ile
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<210> 1448

<211> 276

<212> PRT

<213> Homo sapiens

<400> 1448

Met Val Trp Val Val Leu Leu Ser Leu Leu Cys Tyr Leu Val Leu Phe Leu Cys Arg His Ser Ser His Arg Gly Val Phe Leu Ser Val Thr Ile 25 Leu Ile Tyr Leu Leu Met Gly Glu Met His Met Val Asp Thr Val Thr 45 Trp His Lys Met Arg Gly Ala Gln Met Ile Val Ala Met Lys Ala Val 55 Ser Leu Gly Phe Asp Leu Asp Arg Gly Glu Val Gly Thr Val Pro Ser 75 Pro Val Glu Phe Met Gly Tyr Leu Tyr Phe Val Gly Thr Ile Val Phe 90 Gly Pro Trp Ile Ser Phe His Ser Tyr Leu Gln Ala Val Gln Gly Arg 105 Pro Leu Ser Cys Arg Trp Leu Gln Lys Val Ala Arg Ser Leu Ala Leu 120 125 Ala Leu Leu Cys Leu Val Leu Ser Thr Cys Val Gly Pro Tyr Leu Phe

130 135 140 Pro Tyr Phe Ile Pro Leu Asn Gly Asp Arg Leu Leu Arg Lys Trp Leu 150 155 Arg Ala Tyr Glu Ser Ala Val Ser Phe His Phe Ser Asn Tyr Phe Val 165 170 Gly Phe Leu Ser Glu Ala Thr Ala Thr Leu Ala Gly Ala Gly Phe Thr 185 Glu Glu Lys Asp His Leu Glu Trp Asp Leu Thr Val Ser Lys Pro Leu 200 Asn Val Glu Leu Pro Arg Ser Met Val Glu Val Val Thr Ser Trp Asn 215 220 Leu Pro Met Ser Tyr Trp Leu Asn Asn Tyr Gly Phe Lys Asn Ala Leu 230 235 Arg Leu Gly Thr Leu Leu Gly Cys Ala Gly His Leu Cys Ser Gln Arg 245 250 Pro Ser Lys Leu Leu Lys Phe Pro Pro Gly Trp Gly Pro Cys Cys Pro 265 Gly Phe Leu * 275

<210> 1449

<211> 597

<212> PRT

<213> Homo sapiens

<400> 1449 Met Glu Phe Gly Leu Ser Trp Val Phe Leu Val Ala Ile Leu Lys Gly 10 Val Gln Cys Glu Val Gln Leu Val Glu Ser Gly Gly Leu Val Gln 20 25 Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe 40 Ser Ser Tyr Trp Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu 5.5 Val Trp Val Ser Arg Ile Asn Thr Asp Gly Ser Ser Thr Ser Tyr Ala 70 75 Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala Lys Asn 90 85 Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val 100 105 Tyr Tyr Cys Ala Arg Ala Asp Asn Cys Ser Ser Thr Ser Cys Tyr Lys 120 125 Cys Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly 135 140 Ser Ala Ser Ala Pro Thr Leu Phe Pro Leu Val Ser Cys Glu Asn Ser 150 155 Pro Ser Asp Thr Ser Ser Val Ala Val Gly Cys Leu Ala Gln Asp Phe 170 Leu Pro Asp Ser Ile Thr Phe Ser Trp Lys Tyr Lys Asn Asn Ser Asp 185 Ile Ser Ser Thr Arg Gly Phe Pro Ser Val Leu Arg Gly Gly Lys Tyr 200 Ala Ala Thr Ser Gln Val Leu Leu Pro Ser Lys Asp Val Met Gln Gly 215 220 Thr Asp Glu His Val Val Cys Lys Val Gln His Pro Asn Gly Asn Lys 230 235

```
Glu Lys Asn Val Pro Leu Pro Val Ile Ala Glu Leu Pro Pro Lys Val
               245
                                   250
Ser Val Phe Val Pro Pro Arg Asp Gly Phe Phe Gly Asn Pro Arg Lys
                               265
Ser Lys Leu Ile Cys Gln Ala Thr Gly Phe Ser Pro Arg Gln Ile Gln
                           280
Val Ser Trp Leu Arg Glu Gly Lys Gln Val Gly Ser Gly Val Thr Thr
                       295
Asp Gln Val Gln Ala Glu Ala Lys Glu Ser Gly Pro Thr Thr Tyr Lys
                    310
                                       315
Val Thr Ser Thr Leu Thr Ile Lys Glu Ser Asp Trp Leu Ser Gln Ser
                325
                                   330
Met Phe Thr Cys Arg Val Asp His Arg Gly Leu Thr Phe Gln Gln Asn
           340
                               345
Ala Ser Ser Met Cys Val Pro Asp Gln Asp Thr Ala Ile Arg Val Phe
                           360
Ala Ile Pro Pro Ser Phe Ala Ser Ile Phe Leu Thr Lys Ser Thr Lys
                        375
Leu Thr Cys Leu Val Thr Asp Leu Thr Thr Tyr Asp Ser Val Thr Ile
                    390
                                        395
Ser Trp Thr Arg Gln Asn Gly Glu Ala Val Lys Thr His Thr Asn Ile
                                   410
Ser Glu Ser His Pro Asn Ala Thr Phe Ser Ala Val Gly Glu Ala Ser
                               425
Ile Cys Glu Asp Asp Trp Asn Ser Gly Glu Arg Phe Thr Cys Thr Val
                           440
Thr His Thr Asp Leu Pro Ser Pro Leu Lys Gln Thr Ile Ser Arg Pro
                       455
Lys Gly Val Ala Leu His Arg Pro Asp Val Tyr Leu Leu Pro Pro Ala
                   470
                                       475
Arg Glu Gln Leu Asn Leu Arg Glu Ser Ala Thr Ile Thr Cys Leu Val
               485
                                   490
Thr Gly Phe Ser Pro Ala Asp Val Phe Val Gln Trp Met Gln Arg Gly
                               505
Gln Pro Leu Ser Pro Glu Lys Tyr Val Thr Ser Ala Pro Met Pro Glu
                           520
                                               525
Pro Gln Ala Pro Gly Arg Tyr Phe Ala His Ser Ile Leu Thr Val Ser
                       535
                                           540
Glu Glu Glu Trp Asn Thr Gly Glu Thr Tyr Thr Cys Val Val Ala His
                   550
                                       555
Glu Ala Leu Pro Asn Arg Val Thr Glu Arg Thr Val Asp Lys Ser Thr
               565
                                  570
Gly Lys Pro Thr Leu Tyr Asn Val Ser Leu Val Met Ser Asp Thr Ala
           580
                        585
Gly Thr Cys Tyr *
       595 596
```

<210> 1450 <211> 276 <212> PRT <213> Homo sapiens

```
25
Glu Pro Cys Val Asn Glu Gly Met Cys Val Thr Tyr His Asn Gly Thr
                           40
Gly Tyr Cys Lys Cys Pro Glu Gly Phe Leu Gly Glu Tyr Cys Gln His
                       55
Arg Asp Pro Cys Glu Lys Asn Arg Cys Gln Asn Gly Gly Thr Cys Val
                   70
Ala Gln Ala Met Leu Gly Lys Ala Thr Cys Arg Cys Ala Ser Gly Phe
                                  90
               8.5
Thr Gly Glu Asp Cys Gln Tyr Ser Thr Ser His Pro Cys Phe Val Ser
                              105
Arg Pro Cys Leu Asn Gly Gly Thr Cys His Met Leu Ser Arg Asp Thr
                          120
Tyr Glu Cys Thr Cys Gln Val Gly Phe Thr Gly Lys Glu Cys Gln Trp
                      135
                                         140
Thr Asp Ala Cys Leu Ser His Pro Cys Ala Asn Gly Ser Thr Cys Thr
                  150
                                     155
Thr Val Ala Asn Gln Phe Ser Cys Lys Cys Leu Thr Gly Phe Thr Gly
              165
                                 170
Gln Lys Cys Glu Thr Asp Val Asn Glu Cys Asp Ile Pro Gly His Cys
                             185
                                                 190
Gln His Gly Gly Ile Cys Leu Asn Leu Pro Gly Ser Tyr Gln Cys Gln
                    200
                                     205
Cys Leu Gln Gly Phe Thr Gly Gln Tyr Cys Asp Ser Leu Tyr Val Pro
                     215
                                         220
Cys Ala Pro Ser Pro Cys Val Asn Gly Gly Thr Cys Arg Gln Thr Gly
       230
                                    235
Asp Phe Thr Phe Glu Cys Asn Cys Leu Pro Glu Thr Val Arg Arg Gly
        245 250
Thr Glu Leu Trp Glu Arg Asp Arg Glu Val Trp Asn Gly Lys Glu His
                              265
Asp Glu Asn *
       275
```

<210> 1451

<211> 121

<212> PRT

<213> Homo sapiens

<400> 1451

Met Glu Ser Gly Leu Ser Trp Ile Phe Leu Leu Ala Ile Leu Lys Gly 10 Val Gln Cys Glu Val Gln Leu Val Glu Ser Gly Gly Leu Val Gln 25 Pro Gly Arg Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Arg Phe 40 Asp Glu Tyr Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu 55 Glu Trp Val Gly Gly Ile Ser Trp Asn Arg Asp Ser Ile Ala Tyr Ala 70 Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala Gln Ser 90 Tyr Val Tyr Leu Gln Met Asn Ser Leu Arg His Glu Asp Thr Ala Leu 105 Tyr Tyr Cys Thr Lys Leu Arg Ser Ser 115 120 121

<210> 1452 <211> 48 <212> PRT <213> Homo sapiens

<400> 1452

 Met Glu Arg Gly Asn Ala Leu Val Val Leu Arg Ser Leu Leu Trp Pro

 1
 5
 10
 15

 Gly Leu Thr Phe Tyr His Ala Pro Arg Thr Lys Asn Tyr Gly Tyr Val
 20
 25
 30

 Tyr Val Gly Thr Gly Glu Lys Asn Met Asp Leu Pro Phe Met Leu *
 35
 40
 45
 47

<210> 1453 <211> 123 <212> PRT <213> Homo sapiens

<400> 1453 Met Ile Thr Val Gln Phe Ser Tyr Thr Ala Val Lys Trp Leu Leu Asn Cys Phe Val Leu Ile Leu Tyr Val Ile Leu Ser Ile Leu Phe Gln Val 25 Ser Gln Lys Asn Ser Ser Lys Leu Gly Arg Phe Lys Asn Leu Phe Asn 40 His Lys Glu Cys Ser Lys Leu Leu Phe Asn Arg Asn Gln Ala Gln Thr 55 Leu Glu Leu Thr Ala Asp Arg Ile Arg Phe Gly Leu Phe Pro Glu Trp 75 70 Lys His Phe Ser His Thr Thr Ser Leu Cys Thr Ala Lys Met Leu Ala 85 90 Tyr Pro Leu Trp Phe Pro Ser Phe Ser Leu Ala Ser Gln Arg Asn Leu 105 Pro Pro His Pro Leu Tyr Tyr Ile Phe Tyr * 120 122

<210> 1454 <211> 327 <212> PRT <213> Homo sapiens

```
55
Leu Leu His Gly Phe Pro Thr Ser Ser Tyr Asp Trp Tyr Lys Ile Trp
                  70
                                     75
Glu Gly Leu Thr Leu Arg Phe His Arg Val Ile Ala Leu Asp Phe Leu
                                 90
               85
Gly Phe Gly Phe Ser Asp Lys Pro Arg Pro His His Tyr Ser Ile Phe
          100
                105 110
Glu Gln Ala Ser Ile Val Glu Ala Leu Leu Arg His Leu Gly Leu Gln
           120 `
                                           125
Asn Arg Arg Ile Asn Leu Leu Ser His Asp Tyr Gly Asp Ile Val Ala
                     135
Gln Glu Leu Leu Tyr Arg Tyr Lys Gln Asn Arg Ser Gly Arg Leu Thr
                                    155
Ile Lys Ser Leu Cys Leu Ser Asn Gly Gly Ile Phe Pro Glu Thr His
                     170
Arg Pro Leu Leu Gln Lys Leu Lys Asp Gly Gly Val Leu Ser
                             185
Pro Ile Leu Thr Arg Leu Met Asn Phe Phe Val Phe Ser Arg Gly Leu
                         200
Thr Pro Val Phe Gly Pro Tyr Thr Arg Pro Ser Glu Ser Glu Leu Trp
                      215
Asp Met Trp Ala Gly Ile Arg Asn Asp Gly Asn Leu Val Ile Asp
                  230
                                   235
Ser Leu Leu Gln Tyr Ile Asn Gln Arg Lys Lys Phe Arg Arg Arg Trp
              245
                                 250
Val Gly Ala Leu Ala Ser Val Thr Ile Pro Ile His Phe Ile Tyr Gly
               265
Pro Leu Asp Pro Val Asn Pro Tyr Pro Glu Phe Leu Glu Leu Tyr Arg
                         280
Lys Thr Leu Pro Arg Ser Thr Val Ser Ile Leu Asp Asp His Ile Ser
                  295
His Tyr Pro Gln Leu Glu Asp Pro Met Gly Phe Leu Asn Ala Tyr Met
                  310
                                    315
Gly Phe Ile Asn Ser Phe *
              325 326
```

<210> 1455

<211> 57

<212> PRT

<213> Homo sapiens

<400> 1455

<210> 1456

<211> 48

<212> PRT

<213> Homo sapiens

<210> 1457 <211> 459 <212> PRT <213> Homo sapiens

<400> 1457 Met Ser Asp Leu Leu Ser Val Phe Leu His Leu Leu Leu Leu Phe Lys 10 Leu Val Ala Pro Val Thr Phe Arg His His Arg Tyr Asp Asp Leu Val 20 25 Arg Thr Leu Tyr Lys Val Gln Asn Glu Cys Pro Gly Ile Thr Arg Val 40 Tyr Ser Ile Gly Arg Ser Val Glu Gly Arg His Leu Tyr Val Leu Glu 55 Phe Ser Asp His Pro Gly Ile His Glu Pro Leu Glu Pro Glu Val Lys 70 Tyr Val Gly Asn Met His Gly Asn Glu Ala Leu Gly Arg Glu Leu Met 90 Leu Gln Leu Ser Glu Phe Leu Cys Glu Glu Phe Arg Asn Arg Asn Gln 100 105 Arg Ile Val Gln Leu Ile Gln Asp Thr Arg Ile His Ile Leu Pro Ser 120 Met Asn Pro Asp Gly Tyr Glu Val Ala Ala Ala Gln Gly Pro Asn Lys 135 Pro Gly Tyr Leu Val Gly Arg Asn Asn Ala Asn Gly Val Asp Leu Asn 150 Arg Asn Phe Pro Asp Leu Asn Thr Tyr Ile Tyr Tyr Asn Glu Lys Tyr 170 Gly Gly Pro Asn His His Leu Pro Leu Pro Asp Asn Trp Lys Ser Gln 185 Val Glu Pro Glu Thr Arg Ala Val Ile Arg Trp Met His Ser Phe Asn 200 Phe Val Leu Ser Ala Asn Leu His Gly Gly Ala Val Val Ala Asn Tyr 215 220 Pro Tyr Asp Lys Ser Phe Glu His Arg Val Arg Gly Val Arg Arg Thr 230 235 Ala Ser Thr Pro Thr Pro Asp Asp Lys Leu Phe Gln Lys Leu Ala Lys 245 250 Val Tyr Ser Tyr Ala His Gly Trp Met Phe Gln Gly Trp Asn Cys Gly 260 265 270 Asp Tyr Phe Pro Asp Gly Ile Thr Asn Gly Ala Ser Trp Tyr Ser Leu 280 285 Ser Lys Gly Met Gln Asp Phe Asn Tyr Leu His Thr Asn Cys Phe Glu 295 300 Ile Thr Leu Glu Leu Ser Cys Asp Lys Phe Pro Pro Glu Glu Glu Leu

310 315 Gln Arg Glu Trp Leu Gly Asn Arg Glu Ala Leu Ile Gln Phe Leu Glu 325 330 Gln Val His Gln Gly Ile Lys Gly Met Val Leu Asp Glu Asn Tyr Asn 345 Asn Leu Ala Asn Ala Val Ile Ser Val Ser Gly Ile Asn His Asp Val 360 Thr Ser Gly Asp His Gly Asp Tyr Phe Arg Leu Leu Leu Pro Gly Ile 375 380 Tyr Thr Val Ser Ala Thr Ala Pro Gly Tyr Asp Pro Glu Thr Val Thr 390 395 Val Thr Val Gly Pro Ala Glu Pro Thr Leu Val Asn Phe His Leu Lys 405 410 Arg Ser Ile Pro Gln Val Ser Pro Val Arg Arg Ala Pro Ser Arg Arg 420 425 His Gly Val Arg Ala Lys Val Gln Pro Gln Pro Arg Lys Lys Glu Met 440 Glu Met Arg Gln Leu Gln Arg Gly Pro Ala * 455

<210> 1458 <211> 463 <212> PRT <213> Homo sapiens

<400> 1458 Met Ala Arg Val Leu Gly Ala Pro Val Ala Leu Gly Leu Trp Ser Leu 1.0 Cys Trp Ser Leu Ala Ile Ala Thr Pro Leu Pro Pro Thr Ser Ala His 20 25 Gly Asn Val Ala Glu Gly Glu Thr Lys Pro Asp Pro Asp Val Thr Glu 40 Arg Cys Ser Asp Gly Trp Ser Phe Asp Ala Thr Thr Leu Asp Asp Asn 60 55 Gly Thr Met Leu Phe Phe Lys Gly Glu Phe Val Trp Lys Ser His Lys 70 75 Trp Asp Arg Glu Leu Ile Ser Glu Arg Trp Lys Asn Phe Pro Ser Pro 85 90 Val Asp Ala Ala Phe Arg Gln Gly His Asn Ser Val Phe Leu Ile Lys 105 Gly Asp Lys Val Trp Val Tyr Pro Pro Glu Lys Lys Glu Lys Gly Tyr 120 125 Pro Lys Leu Leu Gln Asp Glu Phe Pro Gly Ile Pro Ser Pro Leu Asp 135 Ala Ala Val Glu Cys His Arg Gly Glu Cys Gln Ala Glu Gly Val Leu Phe Phe Gln Gly Asp Arg Glu Trp Phe Trp Asp Leu Ala Thr Gly Thr 170 Met Lys Glu Arg Ser Trp Pro Ala Val Gly Asn Cys Ser Ser Ala Leu 185 Arg Trp Leu Gly Arg Tyr Tyr Cys Phe Gln Gly Asn Gln Phe Leu Arg 200 205 Phe Asp Pro Val Arg Gly Glu Val Pro Pro Arg Tyr Pro Arg Asp Val 215 220 Arg Asp Tyr Phe Met Pro Cys Pro Gly Arg Gly His Gly His Arg Asn 230 235

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Gly Thr Gly His Gly Asn Ser Thr His His Gly Pro Glu Tyr Met Arg
              245
                   250
Cys Ser Pro His Leu Val Leu Ser Ala Leu Thr Ser Asp Asn His Gly
                             265
          260
Ala Thr Tyr Ala Phe Ser Gly Thr His Tyr Trp Arg Leu Asp Thr Ser
            280
                                             285
Arg Asp Gly Trp His Ser Trp Pro Ile Ala His Gln Trp Pro Gln Gly
                      295
Pro Ser Ala Val Asp Ala Ala Phe Ser Trp Glu Glu Lys Leu Tyr Leu
                  310
                                     315
Val Gln Gly Thr Gln Val Tyr Val Phe Leu Thr Lys Gly Gly Tyr Thr
               325
                                  330
Leu Val Ser Gly Tyr Pro Lys Arg Leu Glu Lys Glu Val Gly Thr Pro
                              345
                                                 350
His Gly Ile Ile Leu Asp Ser Val Asp Ala Ala Phe Ile Cys Pro Gly
                          360
Ser Ser Arg Leu His Ile Met Ala Gly Arg Arg Leu Trp Trp Leu Asp
                       375
                                          380
Leu Lys Ser Gly Ala Gln Ala Thr Trp Thr Glu Leu Pro Trp Pro His
                   390
                                      395
Glu Lys Val Asp Gly Ala Leu Cys Met Glu Lys Ser Leu Gly Pro Asn
               405
                                  410
Ser Cys Ser Ala Asn Gly Pro Gly Leu Tyr Leu Ile His Gly Pro Asn
                              425
                                                 430
Leu Tyr Cys Tyr Ser Asp Val Glu Lys Leu Asn Ala Ala Lys Ala Leu
                          440
Pro Gln Pro Gln Asn Val Thr Ser Leu Leu Gly Cys Thr His *
    450
                       455
                                          460 462
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<210> 1459

<211> 187

<212> PRT

<213> Homo sapiens

<400> 1459

Met Gln Pro Ile Val Ala Lys Ala Leu Val Val Leu Leu Glu Val His 10 Pro Leu Gln Asp Gln Ala Glu Ser Gly Arg Leu Gly His Val His Leu 20 25 Leu Cys Ala Pro Ala Ala Leu Gln His Ala Leu Arg Gly Ile Thr Leu 40 His Asn Gly His His Gln Ala Asp His Leu Pro Asp Leu Met His His 55 60 Glu Ala Leu Ala Leu His Pro Asp His Arg Lys Leu Gln Ala Leu Pro 70 75 His Lys Gly Phe Leu Ala Val His Leu Gln Asp Val Ala Ala Gly Thr 85 90 Gly Ile Leu Arg Pro Leu Leu Arg Gly Glu Ile Val Glu Val Val Arg 105 Ala Leu Val Ala Gly Gln Glu Pro Val Asp Leu Leu Gln Arg Leu Gly 120 Ala Gln Ala Val Gly Leu Ile Leu Asn Val Pro Val Leu Val Arg Lys 135 Gly Lys Arg Gly Gln Gln Val Ala Ile Gly Pro Gly Ile Thr Ser Val 150 155 Leu Gly Val Lys Pro Ala Arg Asp Pro Leu Gln Ser Gln Asn Pro Asn

165 170 175
Val Arg Gly Lys Val Ala Val Asp Leu Phe *
180 185 186

<210> 1460 <211> 223 <212> PRT <213> Homo sapiens

<400> 1460 Met Lys Phe Ala Leu Phe Thr Ser Gly Val Ala Leu Thr Leu Ser Phe 1 5 10 Val Phe Met Tyr Ala Lys Cys Glu Asn Glu Pro Phe Ala Gly Val Ser 20 25 Glu Ser Tyr Asn Gly Thr Gly Glu Leu Gly Asn Leu Ile Ala Pro Cys 35 Asn Ala Asn Cys Asn Cys Ser Arg Ser Tyr Tyr Tyr Pro Val Cys Gly 55 Asp Gly Val Gln Tyr Phe Ser Pro Cys Phe Ala Gly Cys Ser Asn Pro 75 Val Ala His Arg Lys Pro Lys Val Tyr Tyr Asn Cys Ser Cys Ile Glu 85 90 Arg Lys Thr Glu Ile Thr Ser Thr Ala Glu Thr Phe Gly Phe Glu Ala 105 Asn Ala Gly Lys Cys Glu Thr His Cys Ala Lys Leu Ala Ile Phe Leu 120 125 Cys Ile Val Phe Ile Gly Asn Ile Phe Thr Phe Met Ala Arg Ser Pro 140 135 Ile Thr Gly Ala Ile Pro Arg Gly Gly Asn His Arg Gln Arg Pro Pro 155 150 Thr Leu Gly Ile Gln Phe Met Ala Leu Arg Thr Leu Trp Thr Thr Pro 170 165 Trp Pro Ser Lys Thr Gly Cys Pro Ile His Gln Pro Gly Ser Leu Trp 180 185 Glu Lys Leu Gly Trp Arg Pro Leu Lys Thr Leu Arg Arg Pro Lys Pro 205 200 Ser Trp Asn Ala Leu Leu Ala Leu Ala His Pro Arg Ser Phe Gln 220 223 215

<210> 1461 <211> 210 <212> PRT <213> Homo sapiens

Arg Val Val Pro Leu Asn Pro Ala Thr Lys Leu Ser Pro Leu Glu Ser 65 75 Gln Met Ala Leu His Thr Lys Ala Val Glu Ala Gly Met Val Phe Gly His Arg Ala Glu His Lys Asp Pro Arg Ser Val Trp Glu Ser Tyr Trp 105 Leu Leu Gly Ser Pro Trp Ala Glu Val Thr Arg Leu His Pro Arg Arg 120 125 Ala Gln Leu Gly Ser Leu Pro Pro Pro Asp Pro Arg Thr Thr His Arg 135 140 Arg Gly Ala Val Ser Ile Phe Leu Lys Gly Pro Phe Gly Asp Leu Val 150 155 Leu Ser Val Glu Arg Thr Asp Val Ala Leu Ser Ser Gln His Ile Pro 165 170 Gly Ser Gly Arg Pro Gln Leu Lys Gln Cys Gln Gly Pro Gln Gly Ser 180 185 His Leu Asp Arg Pro Thr Ala Cys Asn Ser Ala Leu Leu Arg Arg Gln 200 His * 209

<210> 1462

<211> 56

<212> PRT

<213> Homo sapiens

<400> 1462

 Met Ala Val Arg
 Val Leu Trp Gly Gly Leu Ser Leu Leu Arg Val Leu

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 5
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 5
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 Trp Cys Leu Leu Pro Gln Thr Gly Tyr Val His Pro Asp Glu Phe Phe
 20
 25
 30

 Gln Ser Pro Glu Val Met Ala Gly Lys Thr Pro His Val Trp Leu Arg
 45

 Gln Ala Ala Ala Glu Ser Ala *
 55

<210> 1463

<211> 66

<212> PRT

<213> Homo sapiens

<400> 1463

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<210> 1464
<211> 200
<212> PRT
<213> Homo sapiens
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<400> 1464 Met Val Trp Arg Arg Leu Leu Arg Lys Arg Trp Val Leu Ala Leu Val 10 Phe Gly Leu Ser Leu Val Tyr Phe Leu Ser Ser Thr Phe Lys Gln Glu 20 25 Glu Arg Ala Val Arg Asp Arg Asn Leu Leu Gln Val His Asp His Asn 35 40 Gln Pro Ile Pro Trp Lys Val Gln Phe Asn Leu Gly Asn Ser Ser Arg 55 Pro Ser Asn Gln Cys Arg Asn Ser Ile Gln Gly Lys His Leu Ile Thr 70 75 Asp Glu Leu Gly Tyr Val Cys Glu Arg Lys Asp Leu Leu Val Asn Gly 90 Cys Cys Asn Val Asn Val Pro Ser Thr Lys Gln Tyr Cys Cys Asp Gly 105 110 Cys Trp Pro Asn Gly Cys Cys Ser Ala Tyr Glu Tyr Cys Val Ser Cys 125 120 Cys Leu Gln Pro Asn Lys Gln Leu Leu Leu Glu Arg Phe Leu Asn Arg 135 140 Ala Val Ala Phe Gln Asn Leu Phe Met Ala Val Glu Asp His Phe 150 155 Glu Leu Cys Leu Ala Lys Cys Arg Thr Ser Ser Gln Ser Val Gln His 165 170 Glu Asn Thr Tyr Arg Asp Pro Ile Ala Lys Tyr Cys Tyr Gly Glu Ser 🕟 180 185 Pro Pro Glu Leu Phe Pro Ala * 199

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<210> 1465
<211> 46
<212> PRT
<213> Homo sapiens
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<210> 1466
<211> 56
<212> PRT
<213> Homo sapiens
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<210> 1467 <211> 366 <212> PRT <213> Homo sapiens

<400> 1467 Met Arg Gly Gln Val Val Thr Leu Ile Leu Leu Leu Leu Lys Val 10 Tyr Gln Gly Lys Gly Cys Gln Gly Ser Ala Asp His Val Val Ser Ile Ser Gly Val Pro Leu Gln Leu Gln Pro Asn Ser Ile Gln Thr Lys Val Asp Ser Ile Ala Trp Lys Lys Leu Leu Pro Ser Gln Asn Gly Phe His 55 His Ile Leu Lys Trp Glu Asn Gly Ser Leu Pro Ser Asn Thr Ser Asn 75 Asp Arg Phe Ser Phe Ile Val Lys Asn Leu Ser Leu Leu Ile Lys Ala 90 Ala Gln Gln Asp Ser Gly Leu Tyr Cys Leu Glu Val Thr Ser Ile 105 Ser Gly Lys Val Gln Thr Ala Thr Phe Gln Val Phe Val Phe Asp Lys 120 125 Val Glu Lys Pro Arg Leu Gln Gly Gln Gly Lys Ile Leu Asp Arg Gly 135 140 Arg Cys Gln Val Ala Leu Ser Cys Leu Val Ser Arg Asp Gly Asn Val 150 155 Ser Tyr Ala Trp Tyr Arg Gly Ser Lys Leu Ile Gln Thr Ala Gly Asn 170 165 Leu Thr Tyr Leu Asp Glu Glu Val Asp Ile Asn Gly Thr His Thr Tyr 185 Thr Cys Asn Val Ser Asn Pro Val Ser Trp Glu Ser His Thr Leu Asn 200 205 Leu Thr Gln Asp Cys Gln Asn Ala His Gln Glu Phe Arg Phe Trp Pro 215 220 Phe Leu Val Ile Ile Val Ile Leu Ser Ala Leu Phe Leu Gly Thr Leu 230 235 Ala Cys Phe Cys Val Trp Arg Arg Lys Arg Lys Glu Lys Gln Ser Glu 245 250 Thr Ser Pro Lys Glu Phe Leu Thr Ile Tyr Glu Asp Val Lys Asp Leu 265 Lys Thr Arg Arg Asn His Glu Gln Glu Gln Thr Phe Pro Gly Gly Gly 280 Ser Thr Ile Tyr Ser Met Ile Gln Ser Gln Ser Ser Ala Pro Thr Ser 295 Gln Glu Pro Ala Tyr Thr Leu Tyr Ser Leu Ile Gln Pro Ser Arg Lys

<210> 1468 <211> 57 <212> PRT

<213> Homo sapiens

<210> 1469 <211> 110 <212> PRT <213> Homo sapiens

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<210> 1470 <211> 59 <212> PRT <213> Homo sapiens

<400> 1470

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 Pro
 Val
 Cys
 Ala
 Pro
 Val
 Cys
 Ala
 Pro
 Val
 Ile
 Trp
 Ile
 Glu
 Glu
 Glu
 Glu
 Phe
 Phe
 Pro
 Phe
 Leu
 Lys
 Val
 Ser
 Phe
 Tyr
 Ser

 Leu
 Pro
 Val
 Cys
 Ile
 Glu
 Lys
 Ser
 Ser
 Ile
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 58
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<210> 1471 <211> 123 <212> PRT <213> Homo sapiens

<400> 1471 Met Met His Phe Leu Thr Gly Gly Trp Lys Val Leu Phe Ala Cys Val 10 Pro Pro Thr Glu Tyr Cys His Gly Trp Ala Cys Phe Gly Val Ser Ile 25 Leu Val Ile Gly Leu Leu Thr Ala Leu Ile Gly Asp Leu Ala Ser His Phe Gly Cys Thr Val Gly Leu Lys Asp Ser Val Asn Ala Val Val Phe 55 Val Ala Leu Gly Thr Ser Ile Pro Gly Asn Thr Leu Gly Asp Phe Gly 70 75 Gly Val Gly Ser Gln Met Ser Gln Ala Gly Ala Thr Gln Asp Pro Ala 85 90 Glu Met Arg His Val Arg Gln Gln Gly Gly Gly Ala Ala Gly Pro Val 105 Arg Arg Arg Val His Arg Glu Arg Asp Pro Leu 115 120

<210> 1472 <211> 316 <212> PRT <213> Homo sapiens

 <400> 1472

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115 120 Glu Lys Asn Gly Asp Asn Tyr Asn Trp Phe Phe Leu Ala Leu Pro Thr 130 135 140 Thr Phe Ala Val Ile Glu Asn Leu Lys Tyr Leu Leu Phe Thr Arg Asp 150 155 Ala Ser Gln Pro Phe Tyr Leu Gly His Thr Val Ile Phe Gly Asp Leu 165 170 Glu Tyr Val Thr Val Glu Gly Gly Ile Val Leu Ser Arg Glu Leu Met 190 185 Lys Arg Leu Asn Arg Leu Leu Asp Asn Ser Glu Thr Cys Ala Asp Gln 200 205 Ser Val Ile Trp Lys Leu Ser Glu Asp Lys Gln Leu Ala Ile Cys Leu 215 Lys Tyr Ala Gly Val His Ala Glu Asn Ala Glu Asp Tyr Glu Gly Arg 230 235 Asp Val Phe Asn Thr Lys Pro Ile Ala Gln Leu Ile Glu Glu Ala Leu 245 250 Ser Asn Asn Pro Gln Gln Val Val Glu Gly Cys Cys Ser Asp Met Ala 265 Ile Thr Phe Asn Gly Leu Thr Pro Gln Lys Met Glu Val Met Met Tyr 280 285 Gly Leu Tyr Arg Leu Arg Ala Phe Gly His Tyr Phe Asn Asp Thr Leu 295 Val Phe Leu Pro Pro Val Gly Ser Glu Asn Asp * 310

<210> 1473 <211> 65 <212> PRT

<213> Homo sapiens

<210> 1474 <211> 55 <212> PRT <213> Homo sapiens

Glu Asn Leu Ile Phe Glu Leu Asn Gly Tyr Glu Leu Asn Ser Thr Trp 35 40 45 Phe Gly Trp Leu Asn Thr * 50 54

<210> 1475

<211> 128

<212> PRT

<213> Homo sapiens

<221> misc_feature

<222> (1) ... (128)

<223> Xaa = any amino acid or nothing

<400> 1475

Met Lys Phe Gln Leu Phe Leu Ser Tyr Val Phe Ile Thr Gln Val Phe 5 Ser Arg Pro Phe Gln Ser Asn Leu Gly Ser Leu Thr Pro Ala Ser Ser 25 Gln Ile Pro Leu Gln Leu Pro Lys Ala Leu Cys Val Arg Cys Leu Asn 40 Thr Val Xaa Xaa Xaa Xaa Thr Gly Phe Gly Lys Phe Gln Ile Thr 55 Ile Gln Ser Pro Gly Gly Pro Leu Val Leu Ala Arg Pro Trp Ala Ser 75 Lys Phe Pro Ser Pro Lys Phe Xaa Xaa Xaa Xaa Xaa Pro Lys Met 90 Gly Gly Lys Thr Phe Ala Tyr Gly Arg Ile Asn Pro Thr Arg Pro Ala 105 Lys Asn Xaa Xaa Xaa Xaa Xaa Ser Leu Ala Ser Leu Asn Pro Thr 120 125

<210> 1476

<211> 210

<212> PRT

<213> Homo sapiens

<400> 1476

 Met
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 Asn
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105 100 Leu Leu Gly Ser Pro Trp Ala Glu Val Thr Arg Leu His Pro Arg Arg 120 125 Ala Gln Leu Gly Ser Leu Pro Pro Pro Asp Pro Arg Thr Thr His Arg 135 140 Arg Gly Ala Val Ser Ile Phe Leu Lys Gly Pro Phe Gly Asp Leu Val 150 155 Leu Ser Val Glu Arg Thr Asp Val Ala Leu Ser Ser Gln His Ile Pro 165 170 Gly Ser Gly Arg Pro Gln Leu Lys Gln Cys Gln Gly Pro Gln Gly Ser 185 His Leu Asp Arg Pro Thr Ala Cys Asn Ser Ala Leu Leu Arg Arg Gln 200 209

<210> 1477 <211> 57 <212> PRT <213> Homo sapiens

<210> 1478 <211> 97 <212> PRT <213> Homo sapiens

 Average of the control of the contr

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<210> 1479
<211> 113
<212> PRT
<213> Homo sapiens
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<400> 1479 Met Leu Ser Ile Ser Tyr Phe Ser Asn Ser Leu Met Leu Arg Leu Val 10 Pro Leu Ala Ala Tyr Val Leu Ser Tyr Leu Ile Cys Ser Val Leu Leu 25 His Ile Asn Gln Thr Thr Val Thr Thr Tyr Arg Gly Arg Lys Gln Arg 40 Lys Lys Ile Gln Phe Ala Thr Gly Asn His Gln Ser Ala Gln Ser Tyr 55 Ser Glu Leu Leu Ser Leu Ser Leu Ser Phe Ser Ser Leu Leu Ser Pro 75 70 Val Phe Ser Leu Pro Ser Trp Ser Leu Pro Ser Leu Pro Pro Phe Phe 90 Ser His Ser Pro His Gln Lys Gly Ile Met Met Val Pro Arg Ser Val 105

<210> 1480 <211> 91 <212> PRT <213> Homo sapiens

 Arg Leu
 Ser
 Val
 Cys
 Leu
 Leu
 Leu
 Leu
 Thr
 Leu
 Ala
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<210> 1481 <211> 54 <212> PRT <213> Homo sapiens

20 25 30

Phe Leu Ser Leu Arg Leu Glu Thr Leu Thr Phe Phe Val Leu Trp Leu
35 40 45

Val Pro Tyr Leu Ile *
50 53

<210> 1482 <211> 56 <212> PRT <213> Homo sapiens

<210> 1483 <211> 202 <212> PRT <213> Homo sapiens

<400> 1483 Met Leu Leu Leu Gly Leu Cys Leu Gly Leu Ser Leu Cys Val Gly 10 Ser Gln Glu Glu Ala Gln Ser Trp Gly His Ser Ser Glu Gln Asp Gly 25 Leu Arg Val Pro Arg Gln Val Arg Leu Leu Gln Arg Leu Lys Thr Lys 40 Pro Leu Met Thr Glu Phe Ser Val Lys Ser Thr Ile Ile Ser Arg Tyr 55 Ala Phe Thr Thr Val Ser Cys Arg Met Leu Asn Arg Ala Ser Glu Asp 70 75 Gln Asp Ile Glu Phe Gln Met Gln Ile Pro Ala Ala Ala Phe Ile Thr 90 85 Asn Phe Thr Met Leu Ile Gly Asp Lys Val Tyr Gln Gly Glu Ile Thr 100 105 Glu Arg Glu Lys Lys Ser Gly Asp Arg Val Lys Glu Lys Arg Asn Lys 120 Thr Thr Glu Glu Asn Gly Glu Lys Gly Thr Glu Ile Phe Arg Ala Ser Ala Val Ile Pro Ser Lys Asp Lys Ala Ala Phe Phe Leu Ser Tyr Glu 155 Glu Leu Leu Gln Arg Arg Leu Gly Lys Tyr Glu His Ser Ile Ser Val 170 Arg Pro Gln Gln Leu Ser Gly Arg Leu Ser Val Asp Val Asn Ile Leu 185 Glu Ser Ala Gly Ile Ala Ser Leu Glu Val 200

<210> 1484 <211> 477 <212> PRT <213> Homo sapiens

<400> 1484 Met Pro Gln Leu Ser Leu Ser Trp Leu Gly Leu Gly Gln Val Ala Ala 10 Phe Pro Trp Leu Leu Leu Leu Ala Gly Ala Ser Arg Leu Leu Ala 2.0 25 Gly Phe Leu Ala Trp Thr Tyr Ala Phe Tyr Asp Asn Cys Arg Arg Leu 40 Gln Tyr Phe Pro Gln Pro Pro Lys Gln Lys Trp Phe Trp Gly Gln Pro 55 Gly Pro Pro Ala Ile Ala Pro Lys Asp Asp Leu Ser Ile Arg Phe Leu 75 70 Lys Pro Trp Leu Gly Glu Gly Ile Leu Leu Ser Gly Gly Asp Lys Trp 90 Ser Arg His Arg Arg Met Leu Thr Pro Ala Phe His Phe Asn Ile Leu 105 Lys Ser Tyr Ile Thr Ile Phe Asn Lys Ser Ala Asn Ile Met Leu Asp 120 Lys Trp Gln His Leu Ala Ser Glu Gly Ser Ser Cys Leu Asp Met Phe 135 140 Glu His Ile Ser Leu Met Thr Leu Asp Ser Leu Gln Lys Cys Ile Phe 150 155 Ser Phe Asp Ser His Cys Gln Glu Arg Pro Ser Glu Tyr Ile Ala Thr 165 170 Ile Leu Glu Leu Ser Ala Leu Val Glu Lys Arg Ser Gln His Ile Leu 180 185 Gln His Met Asp Phe Leu Tyr Tyr Leu Ser His Asp Gly Arg Arg Phe 200 205 His Arg Ala Cys Arg Leu Val His Asp Phe Thr Asp Ala Val Ile Arg 220 215 Glu Arg Arg Arg Thr Leu Pro Thr Gln Gly Ile Asp Asp Phe Phe Lys 230 235 Asp Lys Ala Lys Ser Lys Thr Leu Asp Phe Ile Asp Val Leu Leu 250 245 Ser Lys Asp Glu Asp Gly Lys Ala Leu Ser Asp Glu Asp Ile Arg Ala 265 Glu Ala Asp Thr Phe Met Phe Gly Gly His Asp Thr Thr Ala Ser Gly 280 285 Leu Ser Trp Val Leu Tyr Asn Leu Ala Arg His Pro Glu Tyr Gln Glu 295 Arg Cys Arg Gln Glu Val Gln Glu Leu Leu Lys Asp Arg Asp Pro Lys 310 315 Glu Ile Glu Trp Asp Asp Leu Ala Gln Leu Pro Phe Leu Thr Met Cys 330 325 Val Lys Glu Ser Leu Arg Leu His Pro Pro Ala Pro Phe Ile Ser Arg 345 Cys Cys Thr Gln Asp Ile Val Leu Pro Asp Gly Arg Val Ile Pro Lys 360 365 Gly Ile Thr Cys Leu Ile Asp Ile Ile Gly Val His His Asn Pro Thr 380 375 Val Trp Pro Asp Pro Glu Val Tyr Asp Pro Phe Arg Phe Asp Pro Glu

395 390 Asn Ser Lys Gly Arg Ser Pro Leu Ala Phe Ile Pro Phe Ser Ala Gly 410 405 Pro Arg Asn Cys Ile Gly Gln Ala Phe Ala Met Ala Glu Met Lys Val 420 425 Val Leu Ala Leu Met Leu Leu His Phe Arg Phe Leu Pro Asp His Thr 435 440 445 Glu Pro Arg Arg Lys Leu Glu Leu Ile Met Arg Ala Glu Gly Gly Leu 455 460 Trp Leu Arg Val Glu Pro Leu Asn Val Ser Leu Gln 470

<210> 1485 <211> 67 <212> PRT <213> Homo sapiens

<210> 1486 <211> 93 <212> PRT <213> Homo sapiens

<210> 1487 <211> 88 <212> PRT

<213> Homo sapiens

<210> 1488 <211> 268 <212> PRT <213> Homo sapiens

245

His Ser Glu Asp Tyr Ser Lys Val Pro Lys Tyr *

<400> 1488 Met Gly Ser Ala Cys Ile Lys Val Thr Lys Tyr Phe Leu Phe Leu Phe Asn Leu Ile Phe Phe Ile Leu Gly Ala Val Ile Leu Gly Phe Gly Val 25 Trp Ile Leu Ala Asp Lys Ser Ser Phe Ile Ser Val Leu Gln Thr Ser 40 Ser Ser Ser Leu Arg Met Gly Ala Tyr Val Phe Ile Gly Val Gly Ala 55 Val Thr Met Leu Met Gly Phe Leu Gly Cys Ile Gly Ala Val Asn Glu 70 75 Val Arg Cys Leu Leu Gly Leu Tyr Phe Ala Phe Leu Leu Leu Ile Leu 90 85 Ile Ala Gln Val Thr Ala Gly Ala Leu Phe Tyr Phe Asn Met Gly Lys 100 105 Leu Lys Gln Glu Met Gly Gly Ile Val Thr Glu Leu Ile Arg Asp Tyr 120 125 Asn Ser Ser Arg Glu Asp Ser Leu Gln Asp Ala Trp Asp Tyr Val Gln 135 140 Ala Gln Val Lys Cys Cys Gly Trp Val Ser Phe Tyr Asn Trp Thr Asp 150 155 Asn Ala Glu Leu Met Asn Arg Pro Glu Val Thr Tyr Pro Cys Ser Cys 170 165 Glu Val Lys Gly Glu Glu Asp Asn Ser Leu Ser Val Arg Lys Gly Phe 185 Cys Glu Ala Pro Gly Asn Arg Thr Gln Ser Gly Asn His Pro Glu Asp 200 Trp Pro Val Tyr Gln Glu Gly Cys Met Glu Lys Val Gln Ala Trp Leu 215 220 Gln Glu Asn Leu Gly Ile Ile Leu Gly Val Gly Val Gly Val Ala Ile 230 235 Ile Glu Leu Leu Gly Met Val Leu Ser Ile Cys Leu Cys Arg His Val

260 265 267

<210> 1489 <211> 832 <212> PRT

<213> Homo sapiens

<400> 1489 Met Thr Leu Ala Leu Ala Tyr Leu Leu Ala Leu Pro Gln Val Leu Asp 10 Ala Asn Arg Cys Phe Glu Lys Gln Ser Pro Ser Ala Leu Ser Leu Gln 20 25 Leu Ala Ala Tyr Tyr Ser Leu Gln Ile Tyr Ala Arg Leu Ala Pro 40 Cys Phe Arg Asp Lys Cys His Pro Leu Tyr Arg Ala Asp Pro Lys Glu 55 Leu Ile Lys Met Val Thr Arg His Val Thr Arg His Glu His Glu Ala 70 Trp Pro Glu Asp Leu Ile Ser Leu Thr Lys Gln Leu His Cys Tyr Asn 90 Glu Arg Leu Leu Asp Phe Thr Gln Ala Gln Ile Leu Gln Gly Leu Arg 105 Lys Gly Val Asp Val Gln Arg Phe Thr Ala Asp Asp Gln Tyr Lys Arg 120 Glu Thr Ile Leu Gly Leu Ala Glu Thr Leu Glu Glu Ser Val Tyr Ser 135 Ile Ala Ile Ser Leu Ala Gln Arg Tyr Ser Val Ser Arg Trp Glu Val 155 150 Phe Met Thr His Leu Glu Phe Leu Phe Thr Asp Ser Gly Leu Ser Thr 165 170 Leu Glu Ile Glu Asn Arg Ala Gln Asp Leu His Leu Phe Glu Thr Leu 185 Lys Thr Asp Pro Glu Ala Phe His Gln His Met Val Lys Tyr Ile Tyr 200 Pro Thr Ile Gly Gly Phe Asp His Glu Arg Leu Gln Tyr Tyr Phe Thr 215 Leu Leu Glu Asn Cys Gly Cys Ala Asp Leu Gly Asn Cys Ala Ile Lys 230 235 Pro Glu Thr His Ile Arg Leu Leu Lys Lys Phe Lys Val Val Ala Ser 245 250 Gly Leu Asn Tyr Lys Lys Leu Thr Asp Glu Asn Met Ser Pro Leu Glu 260 265 Ala Leu Glu Pro Val Leu Ser Ser Gln Asn Ile Leu Ser Ile Ser Lys 280 285 Leu Val Pro Lys Ile Pro Glu Lys Asp Gly Gln Met Leu Ser Pro Ser 295 300 Ser Leu Tyr Thr Ile Trp Leu Gln Lys Leu Phe Trp Thr Gly Asp Pro 310 315 His Leu Ile Lys Gln Val Pro Gly Ser Ser Pro Glu Trp Leu His Ala 325 330 Tyr Asp Val Cys Met Lys Tyr Phe Asp Arg Leu His Pro Gly Asp Leu 345 Ile Thr Val Val Asp Ala Val Thr Phe Ser Pro Lys Ala Val Thr Lys 360 Leu Ser Val Glu Ala Arg Lys Glu Met Thr Arg Lys Ala Ile Lys Thr

375

Val															
385	Lys	His	Phe	Ile	Glu 390	Lys	Pro	Arg	Lys	Arg 395	Asn	Ser	Glu	Asp	Glu 400
	Gln	Glu	Ala	Lys		Ser	Lys	Val	Thr		Ala	Asp	Thr	Leu	
				405					410					415	
His	Leu	Glu	Lys 420	Ser	Leu	Ala	His	Leu 425	Glu	Thr	Leu	Ser	His 430	Ser	Phe
Ile	Leu	Ser	Leu	Lvs	Asn	Ser	Glu		Glu	Thr	Leu	Gln		Tvr	Ser
		435		4			440					445			
His	Leu 450	Tyr	Asp	Leu	Ser	Arg 455	Ser	Glu	Lys	Glu	Lys 460	Leu	His	Asp	Glu
Ala		Ala	Ile	Cvs	Leu		Glv	Gln	Pro	Leu		Met	Tle	Gln	Gln
465				-1-	470	P	<i>1</i>			475					480
Leu	Leu	Glu	Val	Ala	Val	Gly	Pro	Leu	Asp	Ile	Ser	Pro	Lys	Asp	Ile
				485		-			490				-	495	
Val	Gln	Ser	Ala 500	Ile	Met	Lys	Ile	Ile 505	Ser	Ala	Leu	Ser	Gly 510	Gly	Ser
Ala	Asp	Leu	Gly	Glv	Pro	Ara	Asp		Leu	Lvs	Val	Leu		Glv	Val
		515				_	520					525		_	
Val		Ala	Val	His	Ala		Val	Asp	Lys	GTA		Glu	Leu	Val	Ser
D	530	70	T	T	a1	535	T	7	D	Dl	540	7 J -	7	7	7.1
545	GIU	Asp	Leu	ьeu	550	Trp	ьeu	Arg	Pro	555	Cys	Ala	Asp	Asp	560
	Dro	1757	Arg	Dro		Tla	uic	77-3	T.OU		71 A	T.011	G] 17	Gln	
111	FIU	vaı	Arg	565	Arg	116	1115	vaı	570	GIII	116	neu	GT. À	575	Ser
Phe	His	Leu	Thr		Glu	Asn	Ser	Lvs		Len	Val	Phe	Phe		Thr
		200	580	014	014	1105	001	585			, a1		590		
Glu	Ala	Ile	Leu	Lys	Ala	Ser	Trp		Gln	Arq	Gln	Val		Ile	Ala
		595		-			600			_		605	*		
Asp	Ile	Glu	Asn	G] 11	Glu	Δen	Ara	Tvr	Cvs	Leu	Phe	Met	Glu	Len	Len
E				014				-1-	0,70						Lou
	610					615					620				
	610		His			615					620				
Glu 625	610 Ser	Ser		His	Glu 630	615 Ala	Glu	Phe	Gln	His 635	620 Leu	Val	Leu	Leu	Leu 640
Glu 625	610 Ser	Ser	His	His	Glu 630	615 Ala	Glu	Phe	Gln	His 635	620 Leu	Val	Leu	Leu	Leu 640
Glu 625 Gln	610 Ser Ala	Ser Trp	His Pro Leu	His Pro 645	Glu 630 Met	615 Ala Lys	Glu Ser	Phe Glu Leu	Gln Tyr 650	His 635 Val	620 Leu Ile	Val Thr	Leu Asn Met	Leu Asn 655	Leu 640 Pro
Glu 625 Gln Trp	610 Ser Ala Val	Ser Trp Arg	His Pro Leu 660	His Pro 645 Ala	Glu 630 Met Thr	615 Ala Lys Val	Glu Ser Met	Phe Glu Leu 665	Gln Tyr 650 Thr	His 635 Val Arg	620 Leu Ile Cys	Val Thr Thr	Leu Asn Met 670	Leu Asn 655 Glu	Leu 640 Pro Asn
Glu 625 Gln Trp	610 Ser Ala Val	Ser Trp Arg	His Pro Leu	His Pro 645 Ala	Glu 630 Met Thr	615 Ala Lys Val	Glu Ser Met	Phe Glu Leu 665	Gln Tyr 650 Thr	His 635 Val Arg	620 Leu Ile Cys	Val Thr Thr	Leu Asn Met 670	Leu Asn 655 Glu	Leu 640 Pro Asn
Glu 625 Gln Trp Lys	610 Ser Ala Val Glu	Ser Trp Arg Gly 675	His Pro Leu 660 Leu	His Pro 645 Ala Gly	Glu 630 Met Thr	615 Ala Lys Val Glu	Glu Ser Met Val 680	Phe Glu Leu 665 Leu	Gln Tyr 650 Thr	His 635 Val Arg Met	620 Leu Ile Cys Cys	Val Thr Thr Arg 685	Leu Asn Met 670 Ser	Leu Asn 655 Glu Leu	Leu 640 Pro Asn Tyr
Glu 625 Gln Trp Lys	610 Ser Ala Val Glu	Ser Trp Arg Gly 675	His Pro Leu 660	His Pro 645 Ala Gly	Glu 630 Met Thr	615 Ala Lys Val Glu	Glu Ser Met Val 680	Phe Glu Leu 665 Leu	Gln Tyr 650 Thr	His 635 Val Arg Met	620 Leu Ile Cys Cys	Val Thr Thr Arg 685	Leu Asn Met 670 Ser	Leu Asn 655 Glu Leu	Leu 640 Pro Asn Tyr
Glu 625 Gln Trp Lys Asn	610 Ser Ala Val Glu Thr 690	Ser Trp Arg Gly 675 Lys	His Pro Leu 660 Leu	His Pro 645 Ala Gly Met	Glu 630 Met Thr Asn Leu	615 Ala Lys Val Glu Pro 695	Glu Ser Met Val 680 Ala	Phe Glu Leu 665 Leu	Gln Tyr 650 Thr Lys Gly	His 635 Val Arg Met	620 Leu Ile Cys Cys Lys 700	Val Thr Thr Arg 685 Glu	Leu Asn Met 670 Ser Leu	Leu Asn 655 Glu Leu Cys	Leu 640 Pro Asn Tyr Leu
Glu 625 Gln Trp Lys Asn Leu 705	610 Ser Ala Val Glu Thr 690 Leu	Ser Trp Arg Gly 675 Lys Leu	His Pro Leu 660 Leu Gln Asn	Pro 645 Ala Gly Met	Glu 630 Met Thr Asn Leu Ser 710	615 Ala Lys Val Glu Pro 695 Leu	Glu Ser Met Val 680 Ala Leu	Phe Glu Leu 665 Leu Glu Leu	Gln Tyr 650 Thr Lys Gly Pro	His 635 Val Arg Met Val Ser 715	620 Leu Ile Cys Cys Lys 700 Leu	Val Thr Thr Arg 685 Glu Lys	Leu Asn Met 670 Ser Leu Leu	Leu Asn 655 Glu Leu Cys Leu	Leu 640 Pro Asn Tyr Leu Leu 720
Glu 625 Gln Trp Lys Asn Leu 705	610 Ser Ala Val Glu Thr 690 Leu	Ser Trp Arg Gly 675 Lys Leu	His Pro Leu 660 Leu Gln	Pro 645 Ala Gly Met	Glu 630 Met Thr Asn Leu Ser 710	615 Ala Lys Val Glu Pro 695 Leu	Glu Ser Met Val 680 Ala Leu	Phe Glu Leu 665 Leu Glu Leu	Gln Tyr 650 Thr Lys Gly Pro	His 635 Val Arg Met Val Ser 715	620 Leu Ile Cys Cys Lys 700 Leu	Val Thr Thr Arg 685 Glu Lys	Leu Asn Met 670 Ser Leu Leu	Leu Asn 655 Glu Leu Cys Leu	Leu 640 Pro Asn Tyr Leu Leu 720
Glu 625 Gln Trp Lys Asn Leu 705 Glu	610 Ser Ala Val Glu Thr 690 Leu Ser	Ser Trp Arg Gly 675 Lys Leu Arg	His Pro Leu 660 Leu Gln Asn Asp	His Pro 645 Ala Gly Met Gln Glu 725	Glu 630 Met Thr Asn Leu Ser 710 His	615 Ala Lys Val Glu Pro 695 Leu	Glu Ser Met Val 680 Ala Leu	Phe Glu Leu 665 Leu Glu Leu	Gln Tyr 650 Thr Lys Gly Pro Met 730	His 635 Val Arg Met Val Ser 715 Ala	620 Leu Ile Cys Cys Lys 700 Leu	Val Thr Thr Arg 685 Glu Lys Glu	Leu Asn Met 670 Ser Leu Leu Gln	Leu Asn 655 Glu Leu Cys Leu Ile 735	Leu 640 Pro Asn Tyr Leu 720 Thr
Glu 625 Gln Trp Lys Asn Leu 705 Glu	610 Ser Ala Val Glu Thr 690 Leu Ser	Ser Trp Arg Gly 675 Lys Leu Arg	His Pro Leu 660 Leu Gln Asn	His Pro 645 Ala Gly Met Gln Glu 725	Glu 630 Met Thr Asn Leu Ser 710 His	615 Ala Lys Val Glu Pro 695 Leu	Glu Ser Met Val 680 Ala Leu	Phe Glu Leu 665 Leu Glu Leu	Gln Tyr 650 Thr Lys Gly Pro Met 730	His 635 Val Arg Met Val Ser 715 Ala	620 Leu Ile Cys Cys Lys 700 Leu	Val Thr Thr Arg 685 Glu Lys Glu	Leu Asn Met 670 Ser Leu Leu Gln	Leu Asn 655 Glu Leu Cys Leu Ile 735	Leu 640 Pro Asn Tyr Leu 720 Thr
Glu 625 Gln Trp Lys Asn Leu 705 Glu	610 Ser Ala Val Glu Thr 690 Leu Ser Val	Ser Trp Arg Gly 675 Lys Leu Arg	His Pro Leu 660 Leu Gln Asn Asp Thr	Pro 645 Ala Gly Met Gln Glu 725 Val	Glu 630 Met Thr Asn Leu Ser 710 His	615 Ala Lys Val Glu Pro 695 Leu Leu	Glu Ser Met Val 680 Ala Leu His	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys	His 635 Val Arg Met Val Ser 715 Ala Asp	620 Leu Ile Cys Cys Lys 700 Leu Leu Gln	Val Thr Thr Arg 685 Glu Lys Glu Glu	Leu Asn Met 670 Ser Leu Leu Gln Leu 750	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu	Leu 640 Pro Asn Tyr Leu 720 Thr
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp	Pro 645 Ala Gly Met Gln Glu 725 Val	Glu 630 Met Thr Asn Leu Ser 710 His Asn	615 Ala Lys Val Glu Pro 695 Leu Leu Asp	Glu Ser Met Val 680 Ala Leu His Ser Leu 760	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys	His 635 Val Arg Met Val Ser 715 Ala Asp	620 Leu Ile Cys Cys Lys 700 Leu Leu Gln Val	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755	His Pro Leu 660 Leu Gln Asn Asp Thr 740	Pro 645 Ala Gly Met Gln Glu 725 Val	Glu 630 Met Thr Asn Leu Ser 710 His Asn	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu	Glu Ser Met Val 680 Ala Leu His Ser Leu 760	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys	His 635 Val Arg Met Val Ser 715 Ala Asp	620 Leu Ile Cys Cys Lys 700 Leu Gln Val	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu Pro 770	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755 Arg	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp	Pro 645 Ala Gly Met Gln Glu 725 Val Ala Val	Glu 630 Met Thr Asn Leu Ser 710 His Asn Lys	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu His 775	Glu Ser Met Val 680 Ala Leu His Ser Leu 760 Leu	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val Leu	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys Lys Ala	His 635 Val Arg Met Val Ser 715 Ala Asp Cys	620 Leu Ile Cys Cys Lys 700 Leu Gln Val Leu 780	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765 Gln	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe Arg
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu Pro 770	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755 Arg	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp	Pro 645 Ala Gly Met Gln Glu 725 Val Ala Val	Glu 630 Met Thr Asn Leu Ser 710 His Asn Lys	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu His 775	Glu Ser Met Val 680 Ala Leu His Ser Leu 760 Leu	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val Leu	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys Lys Ala	His 635 Val Arg Met Val Ser 715 Ala Asp Cys	620 Leu Ile Cys Cys Lys 700 Leu Gln Val Leu 780	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765 Gln	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe Arg
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu Tyr Trp 785	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu Pro 770 Asp	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755 Arg	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp	His Pro 645 Ala Gly Met Gln 725 Val Ala Val Glu	Glu 630 Met Thr Asn Leu Ser 710 His Asn Lys Asp Leu 790	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu His 775 Gly	Glu Ser Met Val 680 Ala Leu His Ser Leu 760 Leu Arg	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val Leu His	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys Lys Ala Leu	His 635 Val Arg Met Val Ser 715 Ala Asp Cys Ser Arg 795	620 Leu Ile Cys Cys Lys 700 Leu Gln Val Leu 780 Glu	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765 Gln Ala	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr Gln Gly	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro Gly His	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe Arg Glu 800
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu Tyr Trp 785	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu Pro 770 Asp	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755 Arg	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp Ile Glu	His Pro 645 Ala Gly Met Gln 725 Val Ala Val Glu	Glu 630 Met Thr Asn Leu Ser 710 His Asn Lys Asp Leu 790	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu His 775 Gly	Glu Ser Met Val 680 Ala Leu His Ser Leu 760 Leu Arg	Phe Glu Leu 665 Leu Glu Leu Glu Asn 745 Val Leu His	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys Lys Ala Leu	His 635 Val Arg Met Val Ser 715 Ala Asp Cys Ser Arg 795	620 Leu Ile Cys Cys Lys 700 Leu Gln Val Leu 780 Glu	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765 Gln Ala	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr Gln Gly	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro Gly His	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe Arg Glu 800
Glu 625 Gln Trp Lys Asn Leu 705 Glu Ala Leu Tyr Trp 785 Ala	610 Ser Ala Val Glu Thr 690 Leu Ser Val Leu Pro 770 Asp Glu	Ser Trp Arg Gly 675 Lys Leu Arg Thr Leu 755 Arg Ala Ala	His Pro Leu 660 Leu Gln Asn Asp Thr 740 Asp Ile Glu	His Pro 645 Ala Gly Met Gln 725 Val Ala Val Glu Ser 805	Glu 630 Met Thr Asn Leu Ser 710 His Asn Lys Asp Leu 790 Leu	615 Ala Lys Val Glu Pro 695 Leu Leu Asp Leu His 775 Gly Leu	Glu Ser Met Val 680 Ala Leu His Ser Leu 760 Leu Arg	Phe Glu Leu Glu Leu Glu Asn 745 Val Leu His	Gln Tyr 650 Thr Lys Gly Pro Met 730 Cys Lys Ala Leu Val 810	His 635 Val Arg Met Val Ser 715 Ala Asp Cys Ser Arg 795 Arg	G20 Leu Ile Cys Cys Lys 700 Leu Gln Val Leu 780 Glu Gly	Val Thr Thr Arg 685 Glu Lys Glu Glu Ser 765 Gln Ala Thr	Leu Asn Met 670 Ser Leu Leu Gln Leu 750 Thr Gln Gly His	Leu Asn 655 Glu Leu Cys Leu Ile 735 Leu Pro Gly His Gln 815	Leu 640 Pro Asn Tyr Leu 720 Thr Ser Phe Arg Glu 800

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<400> 1490

<210> 1491 <211> 134 <212> PRT <213> Homo sapiens

<400> 1491 Met Thr Thr Phe Pro Pro Arg Lys Met Val Ala Gln Phe Leu Leu 5 10 Val Ala Gly Asn Val Ala Asn Ile Thr Thr Val Ser Leu Trp Glu Glu 25 Phe Ser Ser Ser Asp Leu Ala Asp Leu Arg Phe Leu Asp Met Ser Gln 40 Asn Gln Phe Gln Tyr Leu Pro Asp Gly Phe Leu Arg Lys Met Pro Ser 55 60 Leu Ser His Leu Asn Leu His Gln Asn Cys Leu Met Thr Leu His Ile 70 . 75 Arg Glu His Glu Pro Pro Gly Ala Leu Thr Glu Leu Asp Leu Ser His 85 90 Asn Gln Leu Ser Glu Leu His Leu Ala Pro Gly Leu Ala Ser Cys Leu 105 Gly Ser Leu Arg Leu Phe Asn Leu Ser Ser Asn Gln Leu Leu Gly Val 115 120 Pro Pro Gly Pro Leu Tyr

<210> 1492 <211> 71 <212> PRT <213> Homo sapiens

134

130

Cys Glu Ser Ile Lys Pro Leu Phe Leu Ile Asn Tyr Pro Val Ser Asn 50 55 60

Lys Ser Leu Leu Ala Thr *

<210> 1493 <211> 78 <212> PRT <213> Homo sapiens

<400> 1493 Met Trp Ile Tyr Phe Trp Thr Leu Asn Ser Val Pro Val Ile Tyr Met 10 Ser Thr Leu Met Ser Ile Pro His Tyr Phe Asp Tyr Cys Cys Phe Ile 30 20 25 Val Ser Asp Ile Met Leu Pro Glu Ile Thr Phe Ser Thr Phe Ile Leu 45 40 35 Leu Leu Met Val Ala Leu Ala Ile Arg Gly Pro Leu His Phe Arg Arg 55 60 His Phe Arg Ile Asn Leu Ser Ile Ala Thr Lys Asn Ala * 75 77

<210> 1494 <211> 121 <212> PRT <213> Homo sapiens

<400> 1494 Met Ala Gly Leu Asn Cys Gly Val Ser Ile Ala Leu Leu Gly Val Leu 10 5 Leu Leu Gly Ala Ala Arg Leu Pro Arg Gly Ala Glu Ala Phe Glu Ile 25 Ala Leu Pro Arg Glu Ser Asn Ile Thr Val Leu Ile Lys Leu Gly Thr 40 Pro Thr Leu Leu Ala Lys Pro Cys Tyr Ile Val Ile Ser Lys Arg His 60 55 Ile Thr Met Leu Ser Ile Lys Ser Gly Glu Arg Ile Val Phe Thr Phe 75 70 Ser Cys Gln Ser Pro Glu Asn His Phe Val Ile Glu Ile Gln Lys Asn 90 85 Ile Asp Cys Met Ser Gly Pro Cys Pro Phe Gly Glu Val Gln Leu Gln 105 Pro Ser Thr Ser Leu Leu Pro Thr Leu

120 121

<210> 1495 <211> 91 <212> PRT <213> Homo sapiens

<210> 1496 <211> 72 <212> PRT

<213> Homo sapiens

<210> 1497 <211> 196 <212> PRT <213> Homo sapiens

<400> 1497 Met Ala Pro Arg Ala Leu Pro Gly Ser Ala Val Leu Ala Ala Val 10 Phe Val Gly Gly Ala Val Ser Ser Pro Leu Val Ala Pro Asp Asn Gly 2.0 25 Ser Ser Arg Thr Leu His Ser Arg Thr Glu Thr Thr Pro Ser Pro Ser 40 Asn Asp Thr Gly Asn Gly His Pro Glu Tyr Ile Ala Tyr Ala Leu Val 55 Pro Val Phe Phe Ile Met Gly Leu Phe Gly Val Leu Ile Cys His Leu Leu Lys Lys Gly Tyr Arg Cys Thr Thr Glu Ala Glu Gln Asp Ile 90 Glu Glu Glu Lys Val Glu Lys Ile Glu Leu Asn Asp Ser Val Asn Glu 105 Asn Ser Asp Thr Val Gly Gln Ile Val His Tyr Ile Met Lys Asn Glu 120

<210> 1498

<211> 75

<212> PRT

<213> Homo sapiens

<400> 1498

 Met
 Trp
 Ser
 Gln
 Ile
 Ala
 Phe
 Val
 Arg
 Ile
 Pro
 Phe
 Cys
 Phe
 Ser
 Leu

 Leu
 Ser
 His
 Ser
 Asn
 Ala
 Trp
 Phe
 Val
 Gln
 Lys
 Ala
 Ala
 Ser
 Gln
 Arg

 Gln
 Ala
 Ser
 Thr
 Ala
 Cys
 His
 Cys
 Pro
 Ala
 Glu
 Ala
 Gly
 Gly

 Glu
 Arg
 Ile
 Thr
 Val
 Ser
 Thr
 Thr
 Thr
 Gly
 Ala
 Gln
 Arg
 Asn
 Ala
 Ala
 Met

 Glu
 Arg
 Ile
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 Val
 Ser
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 Thr
 Thr
 Gly
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 Met

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 Ser
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<210> 1499 <211> 62

<212> PRT

<213> Homo sapiens

<400> 1499

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 Leu
 Met
 Val
 Leu
 Glu
 Ala
 Arg
 Phe
 Val
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 Cys

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 Leu
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 Leu
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<210> 1500

<211> 138

<212> PRT

<213> Homo sapiens

<400> 1500

Met Pro Ile Trp Lys Pro Phe Met Ala Trp Met Ala Ala Trp Ala Leu

10 Ala Val Leu Ser Lys Leu Thr Lys Pro Ile His Leu Leu Trp Met Val 25 Ala Arg Ser Ile Asn Thr Leu Glu Glu Met Ile Leu Pro Lys Gly Thr 40 Asn Ile Cys Val Ser Ser Val Ser Pro Asn Ser Phe Ser Leu Leu Leu 55 60 Leu Gln Glu Gly Arg Arg Leu Glu Asp Ala Val Arg Asp Gly Arg Asp 70 75 Gly Arg Gly Gly Ala His Gly Cys Val Leu Leu Asp Ser Gly Glu Gly 85 90 Arg Met Gln Cys Leu Gly His Ser Arg Ala Leu Ser Trp Val Trp His 105 Lys Ala Ile Gly Ile Asp Glu Phe Pro Gly Gln Gly Ala His Leu Glu 120 Arg Ala Arg His Leu Pro Ser His Trp *

<210> 1501 <211> 82 <212> PRT

<213> Homo sapiens

<210> 1502 <211> 54 <212> PRT <213> Homo sapiens

<210> 1503 <211> 62 <212> PRT <213> Homo sapiens

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<210> 1505 <211> 48 <212> PRT <213> Homo sapiens

<210> 1506 <211> 190 <212> PRT <213> Homo sapiens

<400> 1506 Met Trp Leu Leu Gly Pro Leu Cys Leu Leu Leu Ser Ser Ala Ala Glu

10 Ser Gln Leu Leu Pro Gly Asn Asn Phe Thr Asn Glu Cys Asn Ile Pro 25 Gly Asn Phe Val Cys Ser Asn Gly Arg Cys Ile Pro Gly Ala Trp Gln 40 Cys Asp Gly Leu Pro Asp Cys Phe Asp Lys Ser Asp Glu Lys Glu Cys 55 Pro Lys Ala Lys Ser Lys Cys Gly Pro Thr Phe Pro Cys Ala Ser 70 75 Gly Ile His Cys Ile Ile Gly Arg Phe Arg Cys Asn Gly Phe Glu Asp 90 Cys Pro Asp Gly Ser Asp Glu Glu Asn Cys Thr Ala Asn Pro Leu Leu 105 Cys Ser Thr Ala Arg Tyr His Cys Lys Asn Gly Leu Cys Ile Asp Lys 120 125 Ser Phe Ile Cys Asp Gly Gln Asn Asn Cys Gln Asp Asn Ser Asp Glu 135 Glu Ser Cys Glu Ser Ser Gln Val Phe Arg Pro Gln Val Ser Glu Trp 150 155 Gln Ala Arg Pro Arg Asp Leu Cys Ala Arg Trp Asn Ile Pro Phe Leu 170 Gly Arg Leu Glu Arg Pro Trp Ser Phe Thr Ser Ser Gln Gln 185

<210> 1507 <211> 60 <212> PRT <213> Homo sapiens

<210> 1508 <211> 48 <212> PRT <213> Homo sapiens

<210> 1509 <211> 85 <212> PRT <213> Homo sapiens

 <400> 1509

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 Gly His Ile Ala Ser Ile Leu Ile Pro Leu Leu Leu Leu Leu Leu Leu Leu 20

 Val Leu Ala Ala Gly Val Val Phe Trp Tyr Lys Arg Arg Val Gln Gly 35

 Ala Lys Gly Phe His His Gln Arg Met Thr Asn Gly Ala Met Asn Val 50

 Glu Ile Gly Asn Pro Thr Tyr Lys Met Tyr Glu Gly Gly Glu Pro Asp 65

 Asp Val Gly Gly Leu

<210> 1510 <211> 55 <212> PRT <213> Homo sapiens

<210> 1511 <211> 108 <212> PRT <213> Homo sapiens

85 90 95 Gly Gln Arg Gly Pro Arg Glu Glu Met Arg Gly * 100 105 107

<210> 1512 <211> 119 <212> PRT <213> Homo sapiens

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<400> 1512 Met Val Ala Arg Val Trp Ser Leu Met Arg Phe Leu Ile Lys Gly Ser 1 5 10 Val Ala Gly Gly Ala Val Tyr Leu Val Tyr Asp Gln Glu Leu Leu Gly 25 Pro Ser Asp Lys Ser Gln Ala Ala Leu Gln Lys Ala Gly Glu Val Val 40 45 Pro Pro Ala Met Tyr Gln Phe Ser Gln Tyr Val Cys Gln Gln Thr Gly 55 60 Leu Gln Ile Pro Gln Leu Pro Ala Pro Pro Lys Ile Tyr Phe Pro Ile 75 70 Arg Asp Ser Trp Asn Ala Gly Ile Met Thr Val Met Ser Ala Leu Ser 90 Val Ala Pro Ser Lys Ala Arg Glu Tyr Ser Lys Glu Gly Trp Glu Tyr 100 105 Val Lys Ala Arg Thr Lys * 115 118

<210> 1513 <211> 973 <212> PRT <213> Homo sapiens

<400> 1513 Met Val Lys Ser Lys Trp Gly Leu Ala Leu Ala Ala Val Val Thr Val Leu Ser Ser Leu Leu Met Ser Val Gly Leu Cys Thr Leu Phe Gly Leu 25 Thr Pro Thr Leu Asn Gly Gly Glu Ile Phe Pro Tyr Leu Val Val Val 40 Ile Gly Leu Glu Asn Val Leu Val Leu Thr Lys Ser Val Val Ser Thr 55 Pro Val Asp Leu Glu Val Lys Leu Arg Ile Ala Gln Gly Leu Ser Ser 70 75 Glu Ser Trp Ser Ile Met Lys Asn Met Ala Thr Glu Leu Gly Ile Ile 85 90 Leu Ile Gly Tyr Phe Thr Leu Val Pro Ala Ile Gln Glu Phe Cys Leu 100 105 Phe Ala Val Val Gly Leu Val Ser Asp Phe Phe Leu Gln Met Leu Phe 115 120 Phe Thr Thr Val Leu Ser Ile Asp Ile Arg Arg Met Glu Leu Ala Asp 135 140 Leu Asn Lys Arg Leu Pro Pro Glu Ala Cys Leu Pro Ser Ala Lys Pro 150 155

Val	Gly	Gln	Pro	Thr 165	Arg	Tyr	Glu	Arg	Gln 170		Ala	Val	Arg	Pro 175	Ser
Thr	Pro	His			Thr	Leu	Gln				Phe	Arg			Arg
Len	Pro	Lvs	180	T.e.11	λνα	Val	7727	185	Dho	T.e.ii	בות	Λrα	190	Δνα	T.011
LCu	110	195	Arg	шец	Arg	vai	200	IYI	FILE	БСи	AIA	205	1111	Arg	цец
Ala	Gln 210	Arg	Leu	Ile	Met	Ala 215	Gly	Thr	Val	Val	Trp 220	Ile	Gly	Ile	Leu
Val 225	Tyr	Thr	Asp	Pro	Ala 230	Gly	Leu	Arg	Asn	Tyr 235	Leu	Ala	Ala	Gln	Val 240
Thr	Glu	Gln	Ser	Pro 245	Leu	Gly	Glu	Gly	Ala 250	Leu	Ala	Pro	Met	Pro 255	_
Pro	Ser	Gly	Met 260	Leu	Pro	Pro	Ser	His 265		Asp	Pro	Ala	Phe 270		Ile
Phe	Pro	Pro 275		Ala	Pro	Lys	Leu 280		Glu	Asn	Gln	Thr 285		Pro	Gly
Glu	Ser 290		Glu	Arg	Gly	Gly 295		Ala	Glu	Val	Val 300		Asp	Ser	Pro
Val 305	Pro	Glu	Val	Thr	Trp 310		Pro	Glu	Asp	Glu 315		Leu	Trp	Arg	Lys 320
	Ser	Phe	Arg	His		Pro	Thr	Leu	Phe		Tyr	Tyr	Asn	Ile	
				325					330					335	
	Ala		340	_				345					350		
	Leu	355					360					365			_
Arg	Ser 370	Ala	Trp	Pro	Pro	Pro 375	Gly	Pro	Ile	Pro	Ala 380	Gly	His	Trp	Glu
Ala 385	Gly	Pro	Lys	Gly	Pro 390	Gly	Gly	Val	Gln	Ala 395	His	Gly	Asp	Val	Thr 400
Leu	Tyr	Lys	Val	Ala 405	Ala	Leu	Gly	Leu	Ala 410	Thr	Gly	Ile	Val	Leu 415	Val
Leu	Leu	Leu	Leu 420	Cys	Leu	Tyr	Arg	Val 425	Leu	Cys	Pro	Arg	Asn 430	Tyr	Gly
Gln	Leu	Gly 435	Gly	Gly	Pro	Gly	Arg 440	Arg	Arg	Arg	Gly	Glu 445	Leu	Pro	Cys
Asp	Asp 450	Tyr	Gly	Tyr	Ala	Pro 455	Pro	Glu	Thr	Glu	Ile 460	Val	Pro	Leu	Val
Leu 465	Arg	Gly	His	Leu	Met 470	Asp	Ile	Glu	Cys	Leu 475	Ala	Ser	Asp	Gly	Met 480
Leu	Leu	Val	Ser	Cys 485	Cys	Leu	Ala		His 490	Val	Cys	Val	Ťrp	Asp 495	Ala
Gln	Thr	Gly	Asp 500		Leu	Thr	Arg			Arg	Pro	Gly	Arg 510		Arg
Arg	Asp	Ser 515		Val	Gly	Ser	Gly 520		Glu	Ala	Gln	Glu 525		Trp	Glu
Arg	Leu 530		Asp	Gly	Gly	Lys 535		Gly	Pro	Glu	Glu 540		Gly	Asp	Ser
Pro	Pro	Leu	Arg	His	Arg		Arg	Gly	Pro	Pro		Pro	Ser	Leu	Phe
545		_			550					555					560
Gly	Asp	Gln	Pro	Asp 565	Leu	Thr	Cys	Leu	Ile 570	Asp	Thr	Asn	Phe	Ser 575	Ala
	Pro		580					585				_	590		
Val	Cys	Gly 595	Arg	Ser	Arg	Asp	Ser 600	Pro	Gly	Tyr	Asp	Phe 605	Ser	Cys	Leu
Val	Gln 610	Arg	Val	Tyr	Gln	Glu 615	Glu	Gly	Leu	Ala	Ala 620	Val	Cys	Thr	Pro
Ala	Leu	Arg	Pro	Pro	Ser		Gly	Pro	Val	Leu		Gln	Ala	Pro	Glu

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635
                630
Asp Glu Gly Gly Ser Pro Glu Lys Gly Ser Pro Ser Leu Ala Trp Ala
            645
                              650
Pro Ser Ala Glu Gly Ser Ile Trp Ser Leu Glu Leu Gln Gly Asn Leu
                           665
Ile Val Val Gly Arg Ser Ser Gly Arg Leu Glu Val Trp Asp Ala Ile
                        680
Glu Gly Val Leu Cys Cys Ser Ser Glu Glu Val Ser Ser Gly Ile Thr
                    695
Ala Leu Val Phe Leu Asp Lys Arg Ile Val Ala Ala Arg Leu Asn Gly
                 710
                                   715
Ser Leu Asp Phe Phe Ser Leu Glu Thr His Thr Ala Leu Ser Pro Leu
              725
                               730
Gln Phe Arg Gly Thr Pro Gly Arg Gly Ser Ser Pro Ala Ser Pro Val
                            745
Tyr Ser Ser Ser Asp Thr Val Ala Cys His Leu Thr His Thr Val Pro
                        760
Cys Ala His Gln Lys Pro Ile Thr Ala Leu Lys Ala Ala Ala Gly Arg
                     775
Leu Val Thr Gly Ser Gln Asp His Thr Leu Arg Val Phe Arg Leu Glu
                 790
                                   795
Asp Ser Cys Cys Leu Phe Thr Leu Gln Gly His Ser Gly Ala Ile Thr
             805
                            810
Thr Val Tyr Ile Asp Gln Thr Met Val Leu Ala Ser Gly Gln Asp
       820 825
Gly Ala Ile Cys Leu Trp Asp Val Leu Thr Gly Ser Arg Val Ser His
             840
                               845
Val Phe Ala His Arg Gly Asp Val Thr Ser Leu Thr Cys Thr Thr Ser
                    855
Cys Val Ile Ser Ser Gly Leu Asp Asp Leu Ile Ser Ile Trp Asp Arg
                870
                                  875
Ser Thr Gly Ile Lys Phe Tyr Ser Ile Gln Gln Asp Leu Gly Cys Gly
             885 890
Ala Ser Leu Gly Val Ile Ser Asp Asn Leu Leu Val Thr Gly Gly Gln
         900 905 910
Gly Cys Val Ser Phe Trp Asp Leu Asn Tyr Gly Asp Leu Leu Gln Thr
   915 920
                                        925
Val Tyr Leu Gly Lys Asn Ser Glu Ala Gln Pro Ala Arg Gln Ile Leu
              935
                            940
Val Leu Asp Asn Ala Ala Ile Val Cys Asn Phe Gly Ser Glu Leu Ser
                950
                                  955
Leu Val Tyr Val Pro Ser Val Leu Glu Lys Leu Asp *
                               970
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<210> 1514

<211> 77

<212> PRT

<213> Homo sapiens

<400> 1514

 Met Ile Ser Ser Trp Pro Phe Ser Arg Val Val Arg Phe Trp Phe Leu

 1
 5
 10
 15

 His Gln Met Val Leu Asp Leu Cys Leu Gly Gln Gly Val Pro Gln Gln
 20
 25
 30

 Asn Leu Glu Asn Pro Arg Glu Arg Lys Ser Phe Leu Leu Phe Val Arg
 35
 40
 45

Asn Leu Ile Ile Asp Ser Ser Leu Lys Ile Leu Ser Gln Glu Pro Ser
50 55 60
Asn Leu Trp Gln Arg Ile Pro Lys Met Met Thr Thr *
65 70 75 76

<210> 1515 <211> 148 <212> PRT <213> Homo sapiens

<400> 1515 Met Leu Gly Ser Arg Leu Met Thr Leu Thr Val Cys Ala Gly Ala Leu 10 Ala Arg Gly Arg Gly Thr Gly Thr Cys Glu Thr Arg Gln Glu Gly Lys Gly Gln Asn His Ser Thr Leu Ala Trp Pro His Glu Glu Pro Gly Ala Ser Thr Gly Arg Asp Gly Gly Lys Leu Pro Arg Gly Gln Cys Leu Leu Glu Lys Gly Pro Gly Gly Ala Gly Asp Lys Val Ser Lys Ile Phe Pro 75 Ser Cys Ala Leu Ala Leu Leu Leu Ser Leu Ala Asn Pro Gly Pro Arg 85 90 Gly Pro Arg Glu Phe His Leu Cys Trp Gly Trp Leu Asp Arg Gly Val 105 Thr Gln Glu Ala Val His Val Gly Glu Lys Arg Gly Gly Leu Gly Ser 120 125 Gly Arg Lys Gly Gly Trp Trp Pro Gly Trp Asp Pro Gly Cys Arg Asp 135 Val Ile Thr 145 147

<210> 1516 <211> 274 <212> PRT <213> Homo sapiens

120 115 125 Trp Arg Gly Asp Thr Cys Gln Ser Asp Val Asp Glu Cys Ser Ala Arg 135 140 Arg Gly Gly Cys Pro Gln Arg Cys Val Asn Thr Ala Gly Ser Tyr Trp 150 155 Cys Gln Cys Trp Glu Gly His Ser Leu Ser Ala Asp Gly Thr Leu Cys 170 Val Pro Lys Gly Gly Pro Pro Arg Val Ala Pro Asn Pro Thr Gly Val 185 Asp Ser Ala Met Lys Glu Glu Val Gln Arg Leu Gln Ser Arg Val Asp 200 Leu Leu Glu Glu Lys Leu Gln Leu Val Leu Ala Pro Leu His Ser Leu 215 220 Ala Ser Gln Ala Leu Glu His Gly Leu Pro Asp Pro Gly Ser Leu Leu 230 235 Val His Ser Phe Gln Gln Leu Gly Arg Ile Asp Ser Leu Ser Glu Gln 245 250 Ile Ser Phe Leu Glu Glu Gln Leu Gly Ser Cys Ser Cys Lys Lys Asp 265 Ser *

273

<210> 1517 <211> 246 <212> PRT <213> Homo sapiens

<400> 1517 Met Thr Leu Phe Pro Val Leu Leu Phe Leu Val Ala Gly Leu Leu Pro 1 5 10 Ser Phe Pro Ala Asn Glu Asp Lys Asp Pro Ala Phe Thr Ala Leu Leu 20 25 Thr Thr Gln Thr Gln Val Gln Arg Glu Ile Val Asn Lys His Asn Glu 40 Leu Arg Arg Ala Val Ser Pro Pro Ala Arg Asn Met Leu Lys Met Glu 55 60 Trp Asn Lys Glu Ala Ala Ala Asn Ala Gln Lys Trp Ala Asn Gln Cys 70 75 Asn Tyr Arg His Ser Asn Pro Lys Asp Arg Met Thr Ser Leu Lys Cys 85 90 Gly Glu Asn Leu Tyr Met Ser Ser Ala Ser Ser Ser Trp Ser Gln Ala 105 Ile Gln Ser Trp Phe Asp Glu Tyr Asn Asp Phe Asp Phe Gly Val Gly 120 Pro Lys Thr Pro Asn Ala Val Val Gly His Tyr Thr Gln Val Val Trp 135 Tyr Ser Ser Tyr Leu Val Gly Cys Gly Asn Ala Tyr Cys Pro Asn Gln 155 Lys Val Leu Lys Tyr Tyr Val Cys Gln Tyr Cys Pro Ala Gly Asn 165 170 Trp Ala Asn Arg Leu Tyr Val Pro Tyr Glu Gln Gly Ala Pro Cys Ala 185 Ser Cys Pro Asp Asn Cys Asp Asp Gly Leu Cys Thr Asn Gly Cys Lys 200 205 Tyr Glu Asp Leu Tyr Ser Asn Cys Lys Ser Leu Lys Leu Thr Leu Thr

215

Cys Lys His Gln Leu Val Arg Asp Ser Cys Lys Ala Ser Cys Asn Cys 225 230 235 240 Ser Asn Ser Ile Tyr * 245

<210> 1518

<211> 122

<212> PRT

<213> Homo sapiens

<400> 1518

Met Arg Asn Arg Arg Thr Glu Arg Thr Cys Thr Pro Pro Leu Ala Ser 1.0 Pro Tyr Asn Leu Val Pro His Leu Gln Asn Leu Leu Ala Val Leu Leu 2.0 25 • Met Ile Leu Val Leu Thr Pro Met Val Leu Asn Pro His Lys Leu Tyr 40 Gln Met Met Thr Gln Asn Ile Leu Leu Gln Lys Pro Gln Lys Asn Phe 55 Ile Trp Thr Ala Leu Lys Gly Asn Leu Ser Tyr Pro Arg Asn Leu Leu 70 75 Leu Gln Ser His Leu Ser Leu Leu His Ser Leu Leu Glu Leu 90 Asn Gln Arg Val Cys Leu Leu Pro Arg Ser Leu Ile Asp Pro Gly Lys 105 Arg Leu Lys Lys Pro Met Glu Thr Phe 115 120

<210> 1519

<211> 249

<212> PRT

<213> Homo sapiens

<400> 1519

Met Gly Leu Ser Ile Phe Leu Leu Cys Val Leu Gly Leu Ser Gln 10 Ala Ala Thr Pro Lys Ile Phe Asn Gly Thr Glu Cys Gly Arg Asn Ser 25 Gln Pro Trp Gln Val Gly Leu Phe Glu Gly Thr Ser Leu Arg Cys Gly 40 Gly Val Leu Ile Asp His Arg Trp Val Leu Thr Ala Ala His Cys Ser 55 Gly Ser Arg Tyr Trp Val Arg Leu Gly Glu His Ser Leu Ser Gln Leu 70 75 Asp Trp Thr Glu Gln Ile Arg His Ser Gly Phe Ser Val Thr His Pro 90 Gly Tyr Leu Gly Ala Ser Thr Ser His Glu His Asp Leu Arg Leu Leu 105 Arg Leu Arg Leu Pro Val Arg Val Thr Ser Ser Val Gln Pro Leu Pro 120 Leu Pro Asn Asp Cys Ala Thr Ala Gly Thr Glu Cys His Val Ser Gly 135 140 Trp Gly Ile Thr Asn His Pro Arg Asn Pro Phe Pro Asp Leu Leu Gln

145 150 155 Cys Leu Asn Leu Ser Ile Val Ser His Ala Thr Cys His Gly Val Tyr 170 Pro Gly Arg Ile Thr Ser Asn Met Val Cys Ala Gly Gly Val Pro Gly 185 Gln Asp Ala Cys Gln Gly Asp Ser Gly Gly Pro Leu Val Cys Gly Gly 200 Val Leu Gln Gly Leu Val Ser Trp Gly Ser Val Gly Pro Cys Gly Gln 215 Asp Gly Ile Pro Gly Val Tyr Thr Tyr Ile Cys Lys Tyr Val Asp Trp 230 235 Ile Arg Met Ile Met Arg Asn Asn * 248 245

<210> 1520 <211> 292 <212> PRT <213> Homo sapiens

<400> 1520 Met Leu Val Leu Gln Ile Leu Leu Cys Ile Arg Glu Phe Ile Leu Glu 1 5 1.0 Arg Ser Leu Ile Asn Val Lys Asn Val Ala Lys Ser Leu Ala Val Val 25 Leu Ala Leu Leu Asn Ile Gly Lys Phe Ile Leu Glu Lys Ile Phe Thr 40 45 Asn Ala Lys Tyr Val Leu Asn Leu Leu Leu Val Ser Gln Ile Leu Leu 55 Cys Met Arg Glu Phe Ile Leu Glu Arg Asn Pro Ile Asn Val Lys Asn 70 75 Val Ala Lys Pro Phe Leu Ile Val His Thr Leu Phe Asp Ile Ile Glu 85 90 Phe Ile Leu Glu Lys Asn His Thr Asn Val Lys His Val Ala Asn Leu 105 Leu Val Thr Pro Gln Val Leu Leu Cys Ile Gly Glu Leu Ile Leu Glu 120 Arg Asn Pro Ile His Val Lys Asn Val Ala Lys Pro Leu Val Ile Val 135 Gln Met Leu Phe Ser Ile Gly Glu Phe Ile Leu Ala Arg Asp Pro Thr 150 155 Asn Val Lys Asn Val Ala Lys Pro Ser Thr Ile Gly His Thr Ser Leu 170 His Ile Lys Glu Val Ile Leu Glu Arg Asp Pro Thr Asn Val Lys Asn 185 Val Ala Lys Pro Ser Thr Leu Gly His Thr Ser Leu His Ile Gly Glu 200 Asp Ile Leu Glu Arg Asp Pro Thr Asn Val Met Asn Val Val Lys Pro 215 220 Ser Ala Ile Gly His Thr Ser Leu His Ile Gly Glu Val Ile Val Glu 230 235 Arg Asp Pro Thr Asn Val Lys Asn Val Ala Lys Pro Leu Thr Leu Gly 245 250 His Thr Ser Leu His Ile Arg Glu Val Ile Leu Glu Lys Asn Phe Lys 260 265 Asn Val Lys His Gly Ala Asp Phe Leu Leu Val Thr His Val Leu Leu 280 285

Cys Ile Arg * 290 291

<210> 1521

<211> 129

<212> PRT

<213> Homo sapiens

<400> 1521

120

125

<210> 1522

115

<211> 66

<212> PRT

<213> Homo sapiens

<400> 1522

 Met
 Val
 Val
 Leu
 Pro
 Cys
 Phe
 Ala
 Val
 Leu
 Lys
 Leu
 Phe
 Gly

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 Ser
 Lys
 Leu
 Asp
 Pro
 Pro
 Ser
 Gln
 Ser
 Gly
 Leu
 Asp
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 Ala
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 Gly
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 Arg
 Ala
 Thr
 Asp
 Asp
 Asp
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 Thr
 Ser
 Leu
 Leu
 Leu
 Clu
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<210> 1523

<211> 131

<212> PRT

<213> Homo sapiens

<400> 1523 Met Ile Leu Leu Ala Phe Leu Val Cys Trp Gly Pro Leu Phe Gly Leu 10 Leu Leu Ala Asp Val Phe Gly Ser Asn Leu Trp Ala Gln Glu Tyr Leu 25 Arg Gly Met Asp Trp Ile Leu Ala Leu Ala Val Leu Asn Ser Ala Val 40 Asn Pro Ile Ile Tyr Ser Phe Arg Ser Arg Glu Val Cys Arg Ala Val 55 Leu Ser Phe Leu Cys Cys Gly Cys Leu Arg Leu Gly Met Arg Gly Pro 70 75 Gly Asp Cys Leu Ala Arg Ala Val Glu Ala His Ser Gly Ala Ser Thr 85 90 Thr Asp Ser Ser Leu Arg Pro Arg Asp Ser Phe Arg Gly Ser Arg Ser 105 Leu Ser Phe Arg Met Arg Glu Pro Leu Ser Ser Ile Ser Ser Val Arg Ser Ile * 130

<210> 1524 <211> 52 <212> PRT <213> Homo sapiens

<210> 1525 <211> 246 <212> PRT <213> Homo sapiens

Gly Glu Asn Leu Tyr Met Ser Ser Ala Ser Ser Ser Trp Ser Gln Ala 100 105 Ile Gln Ser Trp Phe Asp Glu Tyr Asn Asp Phe Asp Phe Gly Val Gly 120 125 Pro Lys Thr Pro Asn Ala Val Val Gly His Tyr Thr Gln Val Val Trp 130 135 140 Tyr Ser Ser Tyr Leu Val Gly Cys Gly Asn Ala Tyr Cys Pro Asn Gln 150 155 Lys Val Leu Lys Tyr Tyr Tyr Val Cys Gln Tyr Cys Pro Ala Gly Asn 165 170 Trp Ala Asn Arg Leu Tyr Val Pro Tyr Glu Gln Gly Ala Pro Cys Ala 180 185 Ser Cys Pro Asp Asn Cys Asp Asp Gly Leu Cys Thr Asn Gly Cys Lys 200 Tyr Glu Asp Leu Tyr Ser Asn Cys Lys Ser Leu Lys Leu Thr Leu Thr 215 220 Cys Lys His Gln Leu Val Arg Asp Ser Cys Lys Ala Ser Cys Asn Cys 230 235 Ser Asn Ser Ile Tyr * 245

<210> 1526

<211> 47

<212> PRT

<213> Homo sapiens

<400> 1526

 Met Val Leu Gly Ala Arg Ala Val Ile Ser Phe Cys Ile Leu Ser Ala

 1
 5
 10
 15

 Met Pro Gly Tyr Met Val Val Pro Pro Glu Arg Thr Leu Leu Ala Tyr
 20
 25
 30

 Lys Ser Leu Arg Met Ser Met Ser His Phe Met Met Glu Leu *
 45
 46

<210> 1527

<211> 118

<212> PRT

<213> Homo sapiens

<400> 1527

 Met
 Ser
 Ala
 Arg
 Gly
 Trp
 Pro
 Cys
 Glu
 Ala
 Phe
 Val
 Leu
 Ala
 Gln
 Val

 Cys
 Trp
 Cys
 Trp
 Leu
 Cys
 Val
 Arg
 Gly
 Arg
 Leu
 Cys
 Glu
 Ala
 Leu
 Thr

 Leu
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 Gln
 Val
 Arg
 His
 Gln
 Val
 Cys
 Val
 Pro
 Gly
 Gln
 Pro
 Cys

 Glu
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 Thr
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 Arg
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 His
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 Cys
 Val
 Trp

 Glu
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 Thr
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 Arg
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 Cys
 Val
 Trp

 Gly
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 Trp
 Pro
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 Glu
 Ala
 Leu
 Thr
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100 105 110 Leu Ala Gln Val Arg * 115 117

<210> 1528 <211> 92 <212> PRT <213> Homo sapiens

<210> 1529 <211> 71 <212> PRT <213> Homo sapiens

85

<210> 1530 <211> 85 <212> PRT <213> Homo sapiens

Thr Lys Gly Cys Ile Thr Val Val Gln Gln Ser Gly Ile Leu Thr Glu
35
Leu Lys Gly Gln Gly Ser Phe Leu Tyr Val Leu Leu Cys Leu Asp Ile
50
Thr Leu Leu Val Arg Ser Val Phe Lys Asn Asp Asn Ser Arg Phe Asp
65
70
70
75
80
Phe Gln Ala Asn *

<210> 1531 <211> 60 <212> PRT <213> Homo sapiens

 <400> 1531

 Met Leu Pro Gln Val Phe Leu Gly Phe Thr Lys Val Arg Leu Leu Arg 1

 1
 5
 5
 10
 10
 15
 15

 Leu Arg Asn Pro Trp Gly Cys Val Glu Trp Thr Gly Ala Trp Ser Asp 20
 25
 30
 30
 30
 Arg Arg Trp Asp Gly Ser Gly Val Gly Val Gly Val Gly Leu Asp Pro Thr Cys Pro 35
 45
 45
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<210> 1532 <211> 53 <212> PRT <213> Homo sapiens

<210> 1533 <211> 741 <212> PRT <213> Homo sapiens

52

50

		35					40					45			
Trp	Lys 50	Leu	Val	Ser	Glu	Met 55	Lys	Ala	Glu	Asn	Ile 60	Lys	Ser	Phe	Leu
Arg 65	Ser	Phe	Thr	Lys	Leu 70	Pro	His	Leu	Ala	Gly 75	Thr	Glu	Gln	Asn	Phe 80
Leu	Leu	Ala	Lys	Lys 85	Ile	Gln	Thr	Gln	Trp 90	Lys	Lys	Phe	Gly	Leu 95	Asp
Ser	Ala	Lys	Leu 100	Val	His	Туг	Asp	Val 105	Leu	Leu	Ser	Tyr	Pro 110	Asn	Glu
Thr	Asn	Ala 115	Asn	Tyr	Ile	Ser	Ile 120	Val	Asp	Glu	His	Glu 125	Thr	Glu	Ile
Phe	Lys 130	Thr	Ser	Tyr	Leu	Glu 135	Pro	Pro	Pro	Asp	Gly 140	Tyr	Glu	Asn	Val
Thr 145	Asn	Ile	Val	Pro	Pro 150	Tyr	Asn	Ala	Phe	Ser 155	Ala	Gln	Gly	Met	Pro 160
Glu	Gly	Asp	Leu	Val 165	Tyr	Val	Asn	Tyr	Ala 170	Arg	Thr	Glu	Asp	Phe 175	Phe
	Leu		180					185				_	190		
	Arg	195					200	_		_		205			
	Ala 210					215					220				
225	Ala				230					235					240
	Ala			245	_				250			_		255	_
	Leu		260					265		-			270		_
	Glu	275					280					285			
	Asn 290					295		-	_		300	_			
305	Asp				310					315					320
	Gly			325					330					335	
	Asn		340					345					350		
	Gly	355					360	-				365	-		
	Ser 370					375					380				
385	Gln				390					395					400
	Pro	_	_	405					410	_	=			415	
_	Leu		420					425					430		
	Glu	435				_	440			_		445			-
	Tyr 450					455					460	•			
465	Lys			=	470					475	_	_			480
	Phe			485	_				490					495	
Lys	Asn	Leu	Pro 500	Arg	Ile	Asn	Lys	Leu 505	Gly	Ser	Gly	Ser	Asp 510	Phe	Glu

```
Ala Tyr Phe Gln Arg Leu Gly Ile Ala Ser Gly Arg Ala Arg Tyr Thr
       515
                           520
Lys Asn Lys Lys Thr Asp Lys Tyr Ser Ser Tyr Pro Val Tyr His Thr
                                           540
                       535
Ile Tyr Glu Thr Phe Glu Leu Val Glu Lys Phe Tyr Asp Pro Thr Phe
                                       555
                  550
Lys Lys Gln Leu Ser Val Ala Gln Leu Arg Gly Ala Leu Val Tyr Glu
               565
                                  570
Leu Val Asp Ser Lys Ile Ile Pro Phe Asn Ile Gln Asp Tyr Ala Glu
                               585
Ala Leu Lys Asn Tyr Ala Ala Ser Ile Tyr Asn Leu Ser Lys Lys His
                           600
Asp Gln Gln Leu Thr Asp His Gly Val Ser Phe Asp Ser Leu Phe Ser
                       615
                                           620
Ala Val Lys Asn Phe Ser Glu Ala Ala Ser Asp Phe His Lys Arg Leu
                   630
                                       635
Ile Gln Val Asp Leu Asn Asn Pro Ile Ala Val Arg Met Met Asn Asp
               645
                                   650
Gln Leu Met Leu Glu Arg Ala Phe Ile Asp Pro Leu Gly Leu Pro
                               665
Gly Lys Leu Phe Tyr Arg His Ile Ile Phe Ala Pro Ser Ser His Asn
                           680
                                               685
Lys Tyr Ala Gly Glu Ser Phe Pro Gly Ile Tyr Asp Ala Ile Phe Asp
                       695
                                           700
Ile Glu Asn Lys Ala Asn Ser Arg Leu Ala Trp Lys Glu Val Lys
                    710
                                       715
His Ile Ser Ile Ala Ala Phe Thr Ile Gln Ala Ala Gly Thr Leu
                725
                                   730
Lys Glu Val Leu *
            740
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<210> 1534

<211> 50

<212> PRT

<213> Homo sapiens

<400> 1534

Met Leu Ile Leu Leu His Ile Leu Lys Asn Ile Lys Leu Tyr Leu Val

1 5 10 15

Asn Met Leu Lys Thr Lys Leu Cys Phe Tyr Lys Asp Arg Gly Ser Pro
20 25 30 .

Glu Glu Gly Ile Asp Lys Glu Glu Met Lys Leu Gly Gly Arg Lys Trp
35 40 45

Thr *

49

<210> 1535

<211> 973

<212> PRT

<213> Homo sapiens

<400> 1535

Met Val Lys Ser Lys Trp Gly Leu Ala Leu Ala Val Val Thr Val

,				-					10					7.5	
1 Leu	Ser	Ser	Leu 20	5 Leu	Met	Ser	Val	Gly 25	10 Leu	Cys	Thr	Leu	Phe 30	15 Gly	Leu
Thr	Pro	Thr 35	Leu	Asn	Gly	Gly	Glu 40		Phe	Pro	Tyr	Leu 45		Val	Val
Ile	Gly 50	Leu	Glu	Asn	Val	Leu 55	Val	Leu	Thr	Lys	Ser 60	Val	Val	Ser	Thr
Pro 65	Val	Asp	Leu	Glu	Val 70	Lys	Leu	Arg	Ile	Ala 75	Gln	Gly	Leu	Ser	Ser 80
			Ser	85		_			90				_	95	
			Tyr 100					105					110		
		115	Val	_			120	_				125			
	130		Val			135	_		_		140				
145		-	Arg		150				_	155					160
			Pro	165					170					175	
			Thr 180 Arg					185				_	190		_
		195	Leu		_		200	-				205		_	
	210					215					220				
225			Asp		230					235					240
			Ser	245		_		_	250					255	
		-	Met 260					265		_			270		
		275	Asp			-	280					285			
	290		Glu	_	_	295					300				
305			Val		310	_			_	315					320
			Arg	325					330					335	
			Arg 340					345					350		
		355	Pro	_			360					365			
	370		Trp			375					380				
385			Lys	_	390					395					400
		_	Val	405			_		410					415	
			Leu 420	_			-	425					430		
		435	Gly Gly	_			440					445			
	450		His			455					460				
465	MIG	атХ	nis	neu	470	Asp	TTE	GIU	cys	475	MIA	Ser	Mah	GIY	480

Leu	Leu	Val	Ser	Cys 485	Cys	Leu	Ala	Gly	His 490	Val	Cys	Val	Trp	Asp 495	Ala
Gln	Thr	Gly	Asp 500	Cys	Leu	Thr	Arg	Ile 505	Pro	Arg	Pro	Gly	Arg 510	Gln	Arg
Arg	Asp	Ser 515	Gly	Val	Gly	Ser	Gly 520	Leu	Glu	Ala	Gln	Glu 525	Ser	Trp	Glu
Arg	Leu 530		Asp	Gly	Gly	Lys 535		Gly	Pro	Glu	Glu 540	Pro	Gly	Asp	Ser
Pro 545		Leu	Arg	His	Arg 550		Arg	Gly	Pro	Pro 555		Pro	Ser	Leu	Phe 560
	Asp	Gln	Pro	Asp 565		Thr	Cys	Leu	Ile 570		Thr	Asn	Phe	Ser 575	
Gln	Pro	Arg	Ser 580		Gln	Pro	Thr	Gln 585	-	Glu	Pro	Arg	His 590		Ala
Val	Cys	Gly 595	Arg	Ser	Arg	Asp	Ser 600		Gly	Tyr	Asp	Phe 605		Cys	Leu
.Val			Val	Tyr	Gln	Glu 615		Gly	Leu	Ala	Ala 620		Cys	Thr	Pro
	610 Leu	Arg	Pro	Pro			Gly	Pro	Val	Leu 635		Gln	Ala	Pro	Glu 640
625 Asp	Glu	Gly	Gly		630 Pro	Glu	Lys	Gly	Ser 650		Ser	Leu	Ala	Trp 655	
Pro	Ser	Ala	Glu 660	645 Gly	Ser	Ile	Trp	Ser 665		Glu	Leu	Gln	Gly 670		Leu
Ile	Val	Val 675	Gly	Arg	Ser	Ser	Gly 680		Leu	Glu	Val	Trp 685		Ala	Ile
Glu	Gly 690		Leu	Cys	Cys	Ser 695		Glu	Glu	Val	Ser 700		Gly	Ile	Thr
Ala 705		Val	Phe	Leu	Asp 710		Arg	Ile	Val	Ala 715		Arg	Leu	Asn	Gly 720
	Leu	Asp	Phe	Phe 725		Leu	Glu	Thr	His 730		Ala	Leu	Ser	Pro 735	Leu
Gln	Phe	Arg	Gly 740		Pro	Gly	Arg	Gly 745	Ser	Ser	Pro	Ala	Ser 750	Pro	Val
Tyr	Ser	Ser 755	Ser	Asp	Thr	Val	Ala 760	Cys	His	Leu	Thr	His 765	Thr	Val	Pro
Cys	Ala 770	His	Gln	Lys	Pro	Ile 775	Thr	Ala	Leu	Lys	Ala 780	Ala	Ala	Gly	Arg
Leu 785	Val	Thr	Gly	Ser	Gln 790	Asp	His	Thr	Leu	Arg 795	Val	Phe	Arg	Leu	Glu 800
Asp	Ser	Cys	Cys	Leu 805	Phe	Thr	Leu	Gln	Gly 810	His	Ser	Gly	Ala	Ile 815	Thr
Thr	Val	Tyr	Ile 820	Asp	Gln	Thr	Met	Val 825	Leu	Ala	Ser	Gly	Gly 830	Gln	Asp
Gly	Ala	Ile 835	Cys	Leu	Trp	Asp	Val 840		Thr	Gly	Ser	Arg 845	Val	Ser	His
Val	Phe 850		His	Arg	Gly	Asp 855		Thr	Ser	Leu	Thr 860	Cys	Thr	Thr	Ser
Cys 865		Ile	Ser	Ser	Gly 870	Leu	Asp	Asp	Leu	Ile 875	Ser	Ile	Trp	Asp	Arg 880
Ser	Thr	Gly	Ile	Lys 885		Tyr	Ser	Ile	Gln 890		Asp	Leu	Gly	Cys 895	Gly
			900					905					910		Gln
		915					920					925			
	930					935					940				Leu
Val	Leu	Asp	Asn	Ala	Ala	Ile	Val	Cys	Asn	Phe	Gly	Ser	Glu	Leu	Ser

<210> 1536 <211> 75 <212> PRT

<213> Homo sapiens

<210> 1537 <211> 96 <212> PRT <213> Homo sapiens

<210> 1538 <211> 318 <212> PRT <213> Homo sapiens

Pro Ile Thr Val Thr Gly Ala Gln Val Leu Ser Lys Val Gly Gly Ser 20 25 Val Leu Leu Val Ala Ala Arg Pro Pro Gly Phe Gln Val Arg Glu Ala 40 45 Ile Trp Arg Ser Leu Trp Pro Ser Glu Glu Leu Leu Ala Thr Phe Phe 55 Arg Gly Ser Leu Glu Thr Leu Tyr His Ser Arg Phe Leu Gly Arg Ala 70 Gln Leu His Ser Asn Leu Ser Leu Glu Leu Gly Pro Leu Glu Ser Gly 85 90 Asp Ser Gly Asn Phe Ser Val Leu Met Val Asp Thr Arg Gly Gln Pro 100 105 Trp Thr Gln Thr Leu Gln Leu Lys Val Tyr Asp Ala Val Pro Arg Pro 120 Val Val Gln Val Phe Ile Ala Val Glu Arg Asp Ala Gln Pro Ser Lys 135 140 Thr Cys Gln Val Phe Leu Ser Cys Trp Ala Pro Asn Ile Ser Glu Ile 150 155 -Thr Tyr Ser Trp Arg Arg Glu Thr Thr Met Asp Phe Gly Met Glu Pro 170 His Ser Leu Phe Thr Asp Gly Gln Val Leu Ser Ile Ser Leu Gly Pro 185 Gly Asp Arg Asp Val Ala Tyr Ser Cys Ile Val Ser Asn Pro Val Ser 200 Trp Asp Leu Ala Thr Val Thr Pro Trp Asp Ser Cys His His Glu Ala 215 220 Ala Pro Gly Lys Ala Ser Tyr Lys Asp Val Leu Leu Val Val Pro 230 235 Val Ser Leu Leu Met Leu Val Thr Leu Phe Ser Ala Trp His Trp 245 250 Cys Pro Cys Ser Gly Pro His Leu Arg Ser Lys Gln Leu Trp Met Arg 265 270 Trp Asp Leu Gln Leu Ser Leu His Lys Val Thr Leu Ser Asn Leu Ile 280 285 Ser Thr Val Val Cys Ser Val Val His Gln Gly Leu Val Glu Gln Ile 300 295 His Thr Ala Leu Ile Lys Phe Pro Ser Leu Met Lys Lys 310 315

<210> 1539 <211> 157 <212> PRT <213> Homo sapiens

<400> 1539

 Met Ile Leu Gln Val
 Ser Gly Gly Pro Trp Thr Val Ala Leu Thr Ala

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 Leu Leu Met Val
 Leu Leu Ile Ser Val Val Gln Ser Arg Ala Thr Pro

 20
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 Glu Asn Ser Val Tyr Gln Glu Arg Gln Glu Cys Tyr Ala Phe Asn Gly

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 Thr Gln Arg Val Val Asp Gly Leu Ile Tyr Asn Arg Glu Glu Tyr Val

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 His Phe Asp Ser Ala Val Gly Glu Phe Leu Ala Val Met Glu Leu Gly

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 75
 80

 Arg Pro Ile Gly Glu Tyr Phe Asn Ser Gln Lys Asp Phe Met Glu Arg

| Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution | Solution

<210> 1540 <211> 135 <212> PRT <213> Homo sapiens

<400> 1540 Met Gly Ser Ser Phe Ile Leu Ala Leu Leu Ala Val Leu Gln Gly 1 5 10 Leu Ser Ala Gly Val Leu Leu Glu Gln Ser Arg Ala Glu Val Lys Lys 20 25 Pro Gly Glu Ser Leu Lys Ile Ser Cys Lys Ala Ser Gly Tyr Arg Phe 40 Thr Ser Ala Trp Ile Ala Trp Val Arg Gln Met Pro Gly Lys Gly Leu 55 Glu Trp Met Gly Thr Ile Tyr Pro Ala Asp Ser Glu Val Arg Tyr Ser 70 75 Pro Ser Leu Gln Gly Gln Val Thr Leu Ser Val Asp Glu Ser Ile Ser 90 85 Thr Ala Tyr Leu Gln Trp Asn Ser Leu Arg Ala Ser Asp Thr Ala Thr 105 Tyr Tyr Cys Ala Arg Gln Ile Ile Gly Ala Leu Pro Thr Asp Pro Phe 120 Asp Leu Leu Gly Gln Gly Thr

<210> 1541 <211> 72 <212> PRT <213> Homo sapiens

<210> 1542 <211> 369 <212> PRT <213> Homo sapiens

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<210> 1543 <211> 49 <212> PRT <213> Homo sapiens

<210> 1544 <211> 121 <212> PRT <213> Homo sapiens

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<210> 1545 <211> 70 <212> PRT <213> Homo sapiens

Gln Pro Gly Gln Val * 65 69

<210> 1546

<211> 58

<212> PRT

<213> Homo sapiens

<400> 1546

Cys Cys Val Val Met Ser Pro Leu Pro * 50 55 57

<210> 1547

<211> 65

<212> PRT

<213> Homo sapiens

<400> 1547

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 Trp
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<210> 1548

<211> 78

<212> PRT

<213> Homo sapiens

<400> 1548

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 Leu
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 Phe
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 Leu
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Ser Cys Met Leu Leu Lys Gly Met Ala Cys Leu Leu Thr *

65 70 75 77

<210> 1549

<211> 54

<212> PRT

<213> Homo sapiens

<400> 1549

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 Leu
 Tyr
 Ile
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 Cys
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 Ser
 His
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 Leu
 Val
 Ala
 Pro
 Leu
 Ala

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<212> PRT

<213> Homo sapiens

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 Ala
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 Cys
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 Trp
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<213> Homo sapiens

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 Gln
 Ile
 Asn
 Lys
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 Gly
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 Ser
 Tyr
 Gly
 Pro
 Val
 Ile

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 Leu
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 Ala
 Ser
 Ser
 Val
 Asn
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 Ala
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 Asn
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 Glu
 Ala

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 Leu
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 Gln
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 Gln
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 Glu
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Ala Ser Asn Pro Thr Glu Pro Ala Thr Ile Ile Phe Thr Ala Ala Arg 85 90 Glu Gly Arg Glu Thr Leu Lys Cys Leu Ser His His Val Ala Asp Ala 105 100 Tyr Thr Ser Ser Gln Lys Val Ser Pro Ile Gln Ile Asp Gly Ala Gly 120 125 Arg Thr Trp Gln Asp Ser Asp Thr Val Lys Leu Leu Val Asp Leu Glu 135 140 Leu Ser Tyr Gly Phe Glu Asn Gly Gln Lys Ala Ala Val Wal His His 150 155 Phe Glu Ser Phe Pro Ala Gly Ser Thr Leu Ile Phe Tyr Lys Tyr Cys 165 170 Asp His Glu Asn Ala Ala Phe Lys Asp Val Ala Leu Val Leu Thr Val 185 Leu Glu Glu Glu Thr Leu Glu Ala Ser Val Gly Pro Arg Glu Thr 200 Glu Glu Lys Val Arg Asp Leu Leu Trp Ala Lys Phe Thr Asn Ser * 215 220

<210> 1552

<211> 57

<212> PRT

<213> Homo sapiens

<400> 1552

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 Gln
 Lys
 Phe
 Leu
 Lys
 Pro
 Leu
 Leu
 Ile
 Leu
 His
 Arg
 Leu

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 Lys
 Leu
 Gly
 Ser
 Leu
 Tyr
 Thr
 Pro
 Ser
 Ser
 Val
 Ala
 Arg
 Tyr
 Asp
 Ser

 Ser
 Val
 Asn
 Ser
 Ser
 Val
 Asn
 Ser
 Ser
 Ala
 Tyr
 Glu
 Glu
 Ala

 Lys
 Glu
 Leu
 Met
 Leu
 Ser
 Met
 Asn
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<210> 1553

<211> 241

<212> PRT

<213> Homo sapiens

<400> 1553

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 Ser
 Cys
 Val
 Leu
 Gly
 Gly
 Val
 Ile
 Pro
 Leu
 Gly
 Leu
 Phe
 Leu

 Val
 Cys
 Gly
 Ser
 Gln
 Gly
 Tyr
 Leu
 Leu
 Pro
 Asn
 Val
 Thr
 Leu
 Leu
 Glu

 Glu
 Leu
 Leu
 Ser
 Lys
 Tyr
 Gln
 His
 Ser
 His
 Ser
 Arg
 Val
 Arg

 Arg
 Ala
 Ile
 Pro
 Arg
 Glu
 Asp
 Lys
 Glu
 Glu
 Ile
 Leu
 Met
 Leu
 His
 Asn

 50
 Tyr
 Tyr
 Gln
 Pro
 Gln
 Ala
 Ser
 Asn
 Met
 Glu
 Tyr
 Met

 65
 Tyr
 Tyr
 Glu
 Lys
 Ser
 Ala
 Ala
 Ala
 Ala
 Trp
 Ala
 Ser
 Glu
 Fyr
 But
 Fyr
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100 105 Asn Leu Gly Ala His Trp Gly Arg Tyr Arg Ser Pro Gly Phe His Val 120 Gln Ser Trp Tyr Asp Glu Val Lys Asp Tyr Thr Tyr Pro Tyr Pro Ser 135 140 Glu Cys Asn Pro Trp Cys Pro Glu Arg Cys Ser Gly Pro Met Cys Thr 150 155 His Tyr Thr Gln Ile Val Trp Ala Thr Thr Asn Lys Ile Gly Cys Ala 165 170 175 Val Asn Thr Cys Arg Lys Met Thr Val Trp Gly Glu Val Trp Glu Asn 180 185 190 Ala Val Tyr Phe Val Cys Asn Tyr Ser Pro Lys Gly Asn Trp Ile Gly 200 Glu Ala Pro Tyr Lys Asn Gly Arg Pro Cys Ser Glu Cys Pro Pro Ser 215 Tyr Gly Gly Ser Cys Arg Asn Asn Leu Cys Tyr Arg Glu Glu Thr Tyr 230 235 Thr 241

<210> 1554 <211> 56 <212> PRT

<213> Homo sapiens

<210> 1555 <211> 64 <212> PRT <213> Homo sapiens

<210> 1556

<211> 71 <212> PRT <213> Homo sapiens

<210> 1557 <211> 126 <212> PRT <213> Homo sapiens

<400> 1557 Met Gln Thr His Leu Gly Ala Ser Cys Leu Ser Leu Val Ile Arg Ile 10 Ala Leu Leu Phe Leu Val Gln Arg Asp Gly His Leu His Ser Arg Arg 20 25 Glu Ile Tyr Ala Ile Phe Thr Lys Gly Ser Leu Cys Pro Ala Phe Lys 40 Trp Ala Arg Val Gly Arg Glu Leu Phe Leu His Leu Leu Leu Ser Asn 60 55 Cys His Gln Leu Lys Ile Ile Leu Ile Pro Lys Cys His Ile Leu Gly 70 75 Trp His Ile Leu Ile Pro Phe Thr Ser Lys Ile Trp Asp Ser Tyr Phe 90 85 Ile Val Gln Cys Phe Ser His Phe Thr Thr Leu Ala Asn Val Phe Met 105 Glu Glu Asp Asn Pro Val Ser Glu Leu Gln Val Phe Gln * 120 115

<210> 1558 <211> 135 <212> PRT <213> Homo sapiens

 Phe
 Gln
 Leu
 Pro
 His
 Lys
 Arg
 Glu
 Phe
 Ser
 Glu
 Glu
 Asn
 Pro
 Ala
 Gln
 65
 75
 80
 Asn
 Pro
 Ala
 Ser
 Gly
 Glu
 Asp
 Arg
 Leu
 Trp
 80

 Asn
 Leu
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 Lys
 Val
 Asp
 Ala
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 Arg
 Leu
 Trp
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 Ser
 Val
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 Gln
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 Thr
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 Leu
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 Val
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 Traction
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 Leu
 Cys
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 Asp
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 Met

 Thr
 Asp
 Leu

<210> 1559 <211> 203 <212> PRT <213> Homo sapiens

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<210> 1560 <211> 59 <212> PRT <213> Homo sapiens

Arg Arg Ser Gln Ser Ser Leu Trp Lys Gln Phe Glu Lys Cys Ser Ala 20 25 30

Gly Pro Lys Leu Met Leu Ser Lys Phe Leu Pro Trp Gly Lys Leu Ala 35 40 45

Met Pro Ser Arg Met Ser Asn Phe Ser Pro *
50 55 58

<210> 1561 <211> 50 <212> PRT

<213> Homo sapiens

<210> 1562 <211> 49 <212> PRT <213> Homo sapiens

Ger Thr Asn Val Leu Asn Thr Thr Val Cys Tyr Val Arg Asp Lys Lys 35 40 45 48

<210> 1563 <211> 69 <212> PRT <213> Homo sapiens

50 55 60 His Lys Gln Pro * 65 68

> <210> 1564 <211> 53 <212> PRT <213> Homo sapiens

<210> 1565 <211> 236 <212> PRT <213> Homo sapiens

<400> 1565 Met Pro Arg Arg Gly Leu Ile Leu His Thr Arg Thr His Trp Leu Leu 10 Leu Gly Leu Ala Leu Leu Cys Ser Leu Val Leu Phe Met Tyr Leu Leu 25 Glu Cys Ala Pro Gln Thr Asp Gly Asn Ala Ser Leu Pro Gly Val Val 40 Gly Glu Asn Tyr Gly Lys Glu Tyr Tyr Gln Ala Leu Leu Gln Glu Gln 55 Glu Glu His Tyr Gln Thr Arg Ala Thr Ser Leu Lys Arg Gln Ile Ala 70 75 Gln Leu Lys Gln Glu Leu Gln Glu Met Ser Glu Lys Met Arg Ser Leu 85 90 Gln Glu Arg Arg Asn Val Gly Ala Asn Gly Ile Gly Tyr Gln Ser Asn 100 105 Lys Glu Gln Ala Pro Ser Asp Leu Leu Glu Phe Leu His Ser Gln Ile 120 125 Asp Lys Ala Glu Val Ser Ile Gly Ala Lys Leu Pro Ser Glu Tyr Gly 135 140 Val Ile Pro Phe Glu Ser Phe Thr Leu Met Lys Val Phe Gln Leu Glu 150 155 Met Gly Leu Thr Arg His Pro Glu Glu Lys Pro Val Arg Lys Asp Lys 165 170 Arg Asp Glu Leu Val Glu Val Ile Glu Ala Gly Leu Glu Val Ile Asn 185 Asn Pro Asp Glu Asp Glu Gln Glu Asp Glu Glu Gly Pro Leu Gly 200 205 Glu Lys Leu Ile Phe Asn Glu Asn Asp Phe Val Glu Gly Tyr Tyr Arg

215

Thr Glu Arg Asp Lys Gly Thr Gln Tyr Glu Leu Phe 225 230 235

<210> 1566

<211> 77

<212> PRT

<213> Homo sapiens

<400> 1566

 Met Thr Ala Gly Ile Met Pro Leu Gly Leu Cys Pro Cys Ser Cys Leu

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 Cys Leu His Ser Arg Thr Gly Ala Phe Ser Ala Val His Trp Ser Pro
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 Val Glu Gly Thr Pro Asp Pro Ser Leu Arg Glu Val Ile Ser Lys Gly

 35
 40

Cys Phe Ile Thr Val Phe Pro Gln Asn Asp Pro Ile Asp Thr Val Phe
50 55 60

Ser Gln Cys Pro Leu Thr Phe Glu His Ile Arg Glu * 65 70 75 76

<210> 1567

<211> 104

<212> PRT

<213> Homo sapiens

<400> 1567

 Met
 Leu
 Ile
 Gly
 Leu
 Leu
 Ala
 Trp
 Leu
 Gln
 Thr
 Val
 Pro
 Ala
 His
 Gly

 Cys
 Gln
 Phe
 Leu
 Pro
 Ile
 Thr
 Ser
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 Ala
 Tyr
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 Leu
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 Arg
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 Gln
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Leu Pro His Lys Gln Asp Leu * 100 103

<210> 1568

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1568

Met Val Val Asn Thr Met Ile Tyr Phe Phe Ile Phe Thr Tyr Thr Leu 1 5 10 15 Ala Lys Arg Ala Arg Val His Ile Asn Lys Asn Gly Asn Lys Ala Leu

20 25 Ala Glu Lys Asn Met His Leu Thr Asn His Val Asn Ser * 35 . 40 <210> 1569 <211> 50 <212> PRT <213> Homo sapiens <400> 1569 Met Leu Met Met Asp Thr Leu Trp Pro Ile Leu Leu Gln Thr Leu Lys 10 Val Ile Ser Gln Val Gly His Ala Gly Pro Leu Ala Asn Met Ile His 25 Asp Asn Pro Cys Ile Ile Ala Tyr Arg Ile Thr Leu Arg Leu Val Gly 40 Pro * 49 <210> 1570 <211> 50 <212> PRT <213> Homo sapiens <400> 1570 Met Val Gly Phe Asp Leu Leu Pro Leu Leu Phe Phe Pro Phe Phe 10 Pro Ser Leu Ile Phe Phe Pro Phe Phe Ser Ser Pro Ser Phe 20 25 Gln Phe Leu Pro His Gln Glu Lys Ser Gln His Val Phe Pro Pro Asn 40 Ala * 49 <210> 1571 <211> 50 <212> PRT <213> Homo sapiens <400> 1571 Met Tyr Leu Trp Val Val Arg Trp Lys Trp Cys Leu Gln Lys Leu Gly 5 10 Arg Arg Ile Leu Leu His Ser Leu His Asp Val Phe Ile Ala Asn Met 25 Asp Asp Lys Gly Leu Cys Tyr Arg Gly Leu Arg Ala Pro Ser Phe Leu 40 Leu * 49

<210> 1572 <211> 80 <212> PRT <213> Homo sapiens

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Ser Leu Leu Ser Ser Ser Thr Lys Ser Val Ala Met Pro Thr *
65 70 75 79

<210> 1573 <211> 52 <212> PRT <213> Homo sapiens

(213) Nomo Baptens

<210> 1574 <211> 200 <212> PRT <213> Homo sapiens

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Lys Arg Leu Thr Gly Pro Gly Leu Ser Glu Gly Pro Glu Pro Ser Ile .
                           120
                                             125
Ser Val Met Val Thr Gly Gly Pro Trp His Thr Arg Leu Ser Arg Thr
                       135
                                            140
Cys Leu His Tyr Leu Gly Glu Phe Gly Glu Asp Gln Ile Tyr Glu Ala
                   150
                                        155
His Gln Gln Gly Arg Gly Ala Leu Glu Ala Leu Leu Cys Gly Gly Pro
                165
                                    170
Pro Gly Gly Leu Leu Arg Glu Gly Val Ser His Lys Arg Arg Ala Leu
           180
                                185
Val Leu Asp Ser Thr Leu Leu *
        195
                       199
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<210> 1575
<211> 51
<212> PRT
<213> Homo sapiens
<221> misc_feature
<222> (1) ... (51)

<223> Xaa = any amino acid or nothing

<210> 1576 <211> 124 <212> PRT <213> Homo sapiens

Ser Asp Gly Ala Ser Gly Ser Pro His Thr Gly Glu 115 120 124

> <210> 1577 <211> 860 <212> PRT <213> Homo sapiens

<400> 1577 Met Ala Cys Arg Trp Ser Thr Lys Glu Ser Pro Arg Trp Arg Ser Ala Leu Leu Leu Phe Leu Ala Gly Val Tyr Gly Asn Gly Ala Leu Ala 25 Glu His Ser Glu Asn Val His Ile Ser Gly Val Ser Thr Ala Cys Gly 40 Glu Thr Pro Glu Gln Ile Arg Ala Pro Ser Gly Ile Ile Thr Ser Pro 55 Gly Trp Pro Ser Glu Tyr Pro Ala Lys Ile Asn Cys Ser Trp Phe Ile **7**5 70 Arg Ala Asn Pro Gly Glu Ile Ile Thr Ile Ser Phe Gln Asp Phe Asp 90 Ile Gln Gly Ser Arg Arg Cys Asn Leu Asp Trp Leu Thr Ile Glu Thr 105 Tyr Lys Asn Ile Glu Ser Tyr Arg Ala Cys Gly Ser Thr Ile Pro Pro 120 125 Pro Tyr Ile Ser Ser Gln Asp His Ile Trp Ile Arg Phe His Ser Asp 135 140 Asp Asn Ile Ser Arg Lys Gly Phe Arg Leu Ala Tyr Phe Ser Gly Lys 155 150 Ser Glu Glu Pro Asn Cys Ala Cys Asp Gln Phe Arg Cys Gly Asn Gly 170 165 Lys Cys Ile Pro Glu Ala Trp Lys Cys Asn Asn Met Asp Glu Cys Gly 185 Asp Arg Ser Asp Glu Glu Ile Cys Ala Lys Glu Ala Asn Pro Pro Thr 200 Ala Ala Phe Gln Pro Cys Ala Tyr Asn Gln Phe Gln Cys Leu Ser 215 Arg Phe Thr Lys Val Tyr Thr Cys Leu Pro Glu Ser Leu Lys Cys Asp 230 235 Gly Asn Ile Asp Cys Leu Asp Leu Gly Asp Glu Ile Asp Cys Asp Val 250 245 Pro Thr Cys Gly Gln Trp Leu Lys Tyr Phe Tyr Gly Thr Phe Asn Ser 265 260 Pro Asn Tyr Pro Asp Phe Tyr Pro Pro Gly Ser Asn Cys Thr Trp Leu 280 Ile Asp Thr Gly Asp His Arg Lys Val Ile Leu Arg Phe Thr Asp Phe 300 295 Lys Leu Asp Gly Thr Gly Tyr Gly Asp Tyr Val Lys Ile Tyr Asp Gly 310 315 Leu Glu Glu Asn Pro His Lys Leu Leu Arg Val Leu Thr Ala Phe Asp 330 325 Ser His Ala Pro Leu Thr Val Val Ser Ser Ser Gly Gln Ile Arg Val 340 345 His Phe Cys Ala Asp Lys Val Asn Ala Ala Arg Gly Phe Asn Ala Thr 360 Tyr Gln Val Asp Gly Phe Cys Leu Pro Trp Glu Ile Pro Cys Gly Gly

	370					375					380				
Asn 385	Trp	Gly	Cys	Tyr	Thr 390	Glu	Gln	Gln	Arg	Cys 395		Gly	Tyr	Trp	His 400
Cys	Pro	Asn	Gly	Arg 405	Asp	Glu	Thr	Asn	Cys 410	Thr	Met	Cys	Gln	Lys 415	Glu
Glu	Phe	Pro	Cys 420	Ser	Arg	Asn	Gly	Val 425	Cys	Tyr	Pro	Arg	Ser 430	Asp	Arg
Cys	Asn	Tyr 435	Gln	Asn	His	Cys	Pro 440	Asn	Gly	Ser	Asp	Glu 445	Lys	Asn	Суѕ
Phe	Phe 450	Cys	Gln	Pro	Gly	Asn 455	Phe	His	Cys	Lys	Asn 460	Asn	Arg	Cys	Val
465					470	_			Asp	475					480
				485					Pro 490					495	
			500					505	Leu				510		
		515					520		Arg			525			
	530					535			Ala		540		_		
545					550				Ala	555					560
				565		-			Asn 570					575	
			580					585	Leu				590		
		595		_	_		600		Ile	_		605			
	610			_		615	_		Leu		620				
625					630				Thr	635					640
				645					Val 650					655	_
			660					665	Gly				670		
		675			_		680		Thr			685			
	690					695					700				
705					710				Val	715					720
				725					730					735	Gly
			740					745	Arg				750		
		755			_		760	_	Asn	_		765		_	
	770					775			Pro		780				
785					790				Pro	795					800
		_		805		_			Tyr 810					815	-
			820					825	Сув				830		
uis	ınr	835	GID	тте	PTO	Asp	1hr 840	cys	Leu	ъц	val	845	ьeu	пàs	ASN

Glu Thr Ser Asp Asp Glu Ala Leu Leu Cys * 850 859

<210> 1578 <211> 58 <212> PRT <213> Homo sapiens

<210> 1579 <211> 572 <212> PRT <213> Homo sapiens

<400> 1579 Met Arg Arg Arg Ser Arg Met Leu Leu Cys Phe Ala Phe Leu Trp Val Leu Gly Ile Ala Tyr Tyr Met Tyr Ser Gly Gly Gly Ser Ala Leu Ala Gly Gly Ala Gly Gly Ala Gly Arg Lys Glu Asp Trp Asn Glu Ile Asp Pro Ile Lys Lys Lys Asp Leu His His Ser Asn Gly Glu Glu Lys 55 Ala Gln Ser Met Glu Thr Leu Pro Pro Gly Lys Val Arg Trp Pro Asp 70 Phe Asn Gln Glu Ala Tyr Val Gly Gly Thr Met Val Arg Ser Gly Gln 90 85 Asp Pro Tyr Ala Arg Asn Lys Phe Asn Gln Val Glu Ser Asp Lys Leu 105 Arg Met Asp Arg Ala Ile Pro Asp Thr Arg His Asp Gln Cys Gln Arg 120 Lys Gln Trp Arg Val Asp Leu Pro Ala Thr Ser Val Val Ile Thr Phe 140 135 His Asn Glu Ala Arg Ser Ala Leu Leu Arg Thr Val Val Ser Val Leu 155 150 Lys Lys Ser Pro Pro His Leu Ile Lys Glu Ile Ile Leu Val Asp Asp 170 Tyr Ser Asn Asp Pro Glu Asp Gly Ala Leu Leu Gly Lys Ile Glu Lys 185 Val Arg Val Leu Arg Asn Asp Arg Arg Glu Gly Leu Met Arg Ser Arg . 205 200 Val Arg Gly Ala Asp Ala Ala Gln Ala Lys Val Leu Thr Phe Leu Asp 220 215 Ser His Cys Glu Cys Asn Glu His Trp Leu Glu Pro Leu Leu Glu Arg

```
235
225
                  230
Val Ala Glu Asp Arg Thr Arg Val Val Ser Pro Ile Ile Asp Val Ile
                      250
            245
Asn Met Asp Asn Phe Gln Tyr Val Gly Ala Ser Ala Asp Leu Lys Gly
                             265 270
Gly Phe Asp Trp Asn Leu Val Phe Lys Trp Asp Tyr Met Thr Pro Glu
                         280
Gln Arg Arg Ser Arg Gln Gly Asn Pro Val Ala Pro Ile Lys Thr Pro
                      295
Met Ile Ala Gly Gly Leu Phe Val Met Asp Lys Phe Tyr Phe Glu Glu
                                    315
                  310
Leu Gly Lys Tyr Asp Met Met Asp Val Trp Gly Gly Glu Asn Leu
              325
                                 330
Glu Ile Ser Phe Arg Val Trp Gln Cys Gly Gly Ser Leu Glu Ile Ile
                              345
Pro Cys Ser Arg Val Gly His Val Phe Arg Lys Gln His Pro Tyr Thr
                          360
Phe Pro Gly Gly Ser Gly Thr Val Phe Ala Arg Asn Thr Arg Arg Ala
                      375
Ala Glu Val Trp Met Asp Glu Tyr Lys Asn Phe Tyr Tyr Ala Ala Val
                  390
                                     395
Pro Ser Ala Arg Asn Val Pro Tyr Gly Asn Ile Gln Ser Arg Leu Glu
                                 410 415
Leu Arg Lys Lys Leu Ser Cys Lys Pro Phe Lys Trp Tyr Leu Glu Asn
        420
                             425
Val Tyr Pro Glu Leu Arq Val Pro Asp His Gln Asp Ile Ala Phe Gly
                       440
Ala Leu Gln Gln Gly Thr Asn Cys Leu Asp Thr Leu Gly His Phe Ala
                      455
Asp Gly Val Val Gly Val Tyr Glu Cys His Asn Ala Gly Gly Asn Gln
                 470
                                     475
Glu Trp Ala Leu Thr Lys Glu Lys Ser Val Lys His Met Asp Leu Cys
              485
                             490
Leu Thr Val Val Asp Arg Ala Pro Gly Ser Leu Ile Lys Leu Gln Gly
                             505
Cys Arg Glu Asn Asp Ser Arg Gln Lys Trp Glu Gln Ile Glu Gly Asn
                         520
Ser Lys Leu Arg His Val Gly Ser Asn Leu Cys Leu Asp Ser Arg Thr
                     535
                                       540
Ala Lys Ser Gly Gly Leu Ser Val Glu Val Cys Gly Pro Ala Leu Ser
                 550
                                    555
Gln Gln Trp Lys Phe Thr Leu Asn Leu Gln Gln *
               565
                                 570 571
```

<210> 1580

<211> 77

<212> PRT

<213> Homo sapiens

<400> 1580

 Met Glu Arg Pro Leu Cys Ser His Leu Cys Ser Cys Leu Ala Met Leu

 1
 5
 10
 15

 Ala Leu Leu Ser Pro Leu Ser Leu Ala Gln Tyr Asp Ser Trp Pro His
 20
 25
 30

 Tyr Pro Glu Tyr Phe Gln Gln Pro Ala Pro Glu Tyr His Gln Pro Gln
 35
 40
 45

Ala Pro Ala Asn Val Ala Lys Ile Gln Leu Arg Leu Ala Gly Gln Lys
50 55 60

Arg Lys His Ser Glu Gly Pro Gly Gly Gly Val Leu *
65 70 75 76

<210> 1581 <211> 494 <212> PRT <213> Homo sapiens

<400> 1581 Met Gly Ser Leu Gln Pro Leu Ala Thr Leu Tyr Leu Leu Gly Met Leu 10 Val Ala Ser Cys Leu Gly Arg Leu Ser Trp Tyr Asp Pro Asp Phe Gln 25 20 Ala Arg Leu Thr Arg Ser Asn Ser Lys Cys Gln Gly Gln Leu Glu Val 40 Tyr Leu Lys Asp Gly Trp His Met Val Cys Ser Gln Ser Trp Gly Arg 55 Ser Ser Lys Gln Trp Glu Asp Pro Ser Gln Ala Ser Lys Val Cys Gln 75 70 Arg Leu Asn Cys Gly Val Pro Leu Ser Leu Gly Pro Phe Leu Val Thr 85 90 Tyr Thr Pro Gln Ser Ser Ile Ile Cys Tyr Gly Gln Leu Gly Ser Phe 105 Ser Asn Cys Ser His Ser Arg Asn Asp Met Cys His Ser Leu Gly Leu 125 120 Thr Cys Leu Glu Pro Gln Lys Thr Thr Pro Pro Thr Thr Arg Pro Pro 140 135 Pro Thr Thr Pro Glu Pro Thr Ala Pro Pro Arg Leu Gln Leu Val 155 Ala Gln Ser Gly Gly Gln His Cys Ala Gly Val Val Glu Phe Tyr Ser 170 175 Gly Ser Leu Gly Gly Thr Ile Ser Tyr Glu Ala Gln Asp Lys Thr Gln 190 185 Asp Leu Glu Asn Phe Leu Cys Asn Asn Leu Gln Cys Gly Ser Phe Leu 205 200 Lys His Leu Pro Glu Thr Glu Ala Gly Arg Ala Gln Asp Pro Gly Glu 220 215 Pro Arg Glu His Gln Pro Leu Pro Ile Gln Trp Lys Ile Gln Asn Ser 230 235 Ser Cys Thr Ser Leu Glu His Cys Phe Arg Lys Ile Lys Pro Gln Lys 250 245 Ser Gly Arg Val Leu Ala Leu Leu Cys Ser Gly Phe Gln Pro Lys Val 265 Gln Ser Arg Leu Val Gly Gly Ser Ser Ile Cys Glu Gly Thr Val Glu 280 Val Arg Gln Gly Ala Gln Trp Ala Ala Leu Cys Asp Ser Ser Ser Ala 295 Arg Ser Ser Leu Arg Trp Glu Glu Val Cys Arg Glu Gln Gln Cys Gly 315 310 Ser Val Asn Ser Tyr Arg Val Leu Asp Ala Gly Asp Pro Thr Ser Arg 330 325 Gly Leu Phe Cys Pro His Gln Lys Leu Ser Gln Cys His Glu Leu Trp 345 Glu Arg Asn Ser Tyr Cys Lys Lys Val Phe Val Thr Cys Gln Asp Pro

355 360 365 Asn Pro Ala Gly Leu Ala Ala Gly Thr Val Ala Ser Ile Ile Leu Ala 375 380 Leu Val Leu Leu Val Leu Leu Val Val Cys Gly Pro Leu Ala Tyr 390 395 Lys Lys Leu Val Lys Lys Phe Arg Gln Lys Lys Gln Arg Gln Trp Ile 405 410 Gly Pro Thr Gly Met Asn Gln Asn Met Ser Phe His Arg Asn His Thr 425 Ala Thr Val Arg Ser His Ala Glu Asn Pro Thr Ala Ser His Val Asp 440 Asn Glu Tyr Ser Gln Pro Pro Arg Asn Ser Arg Leu Ser Ala Tyr Pro 455 460 Ala Leu Glu Gly Ala Leu His Arg Ser Ser Met Gln Pro Asp Asn Ser 470 475 Ser Asp Ser Asp Tyr Asp Leu His Gly Ala Gln Arg Leu * 485 490 493

<210> 1582 <211> 329 <212> PRT <213> Homo sapiens

<400> 1582 Met Gln Gly Leu Cys Ile Ser Val Ala Val Phe Leu His Tyr Phe Leu 10 Leu Val Ser Phe Thr Trp Met Gly Leu Glu Ala Phe His Met Tyr Leu 25 Ala Leu Val Lys Val Phe Asn Thr Tyr Ile Arg Lys Tyr Ile Leu Lys 40 4.5 Phe Cys Ile Val Gly Trp Gly Val Pro Ala Val Val Thr Ile Ile 55 Leu Thr Ile Ser Pro Asp Asn Tyr Gly Leu Gly Ser Tyr Gly Lys Phe 70 75 Pro Asn Gly Ser Pro Asp Asp Phe Cys Trp Ile Asn Asn Asn Ala Val 85 90 Phe Tyr Ile Thr Val Val Gly Tyr Phe Cys Val Ile Phe Leu Leu Asn 100 105 Val Ser Met Phe Ile Val Val Leu Val Gln Leu Cys Arg Ile Lys Lys 120 125 Lys Lys Gln Leu Gly Ala Gln Arg Lys Thr Ser Ile Gln Asp Leu Arg 135 140 Ser Ile Ala Gly Leu Thr Phe Leu Leu Gly Ile Thr Trp Gly Phe Ala 150 155 Phe Phe Ala Trp Gly Pro Val Asn Val Thr Phe Met Tyr Leu Phe Ala 170 Ile Phe Asn Thr Leu Gln Gly Phe Phe Ile Phe Ile Phe Tyr Cys Val 185 Ala Lys Glu Asn Val Arg Lys Gln Trp Arg Arg Tyr Leu Cys Cys Gly 200 Lys Leu Arg Leu Ala Glu Asn Ser Asp Trp Ser Lys Thr Ala Thr Asn 215 220 Gly Leu Lys Lys Gln Thr Val Asn Gln Gly Val Ser Ser Ser Asn 230 235 Ser Leu Gln Ser Ser Ser Asn Ser Thr Asn Ser Thr Thr Leu Leu Val 250

<210> 1583 <211> 49 <212> PRT <213> Homo sapiens

<210> 1584 <211> 671 <212> PRT <213> Homo sapiens

<400> 1584 Met Ile Ala Ser Cys Leu Cys Tyr Leu Leu Leu Pro Ala Thr Arg Leu Phe Arg Ala Leu Ser Asp Ala Phe Phe Thr Cys Arg Lys Asn Val Leu 25 Leu Ala Asn Ser Ser Pro Gln Val Glu Gly Asp Phe Ala Met Ala 40 Pro Arg Gly Pro Glu Gln Glu Cys Glu Gly Leu Leu Gln Gln Trp 55 Arg Glu Glu Gly Leu Ser Gln Val Leu Ser Thr Ala Ser Glu Gly Pro 70 75 Leu Ile Asp Lys Gly Leu Ala Gln Ser Ser Leu Ala Leu Leu Met Asp 85 90 Asn Pro Gly Glu Asn Ala Ala Ser Glu Asp Arg Trp Ser Ser Arg 105 110 Gln Leu Ser Asp Leu Arg Ala Ala Glu Asn Leu Asp Glu Pro Phe Pro 120 125 Glu Met Leu Gly Glu Glu Pro Leu Leu Glu Val Glu Gly Val Glu Gly 135 140 Ser Met Trp Ala Ala Ile Pro Met Gln Ser Glu Pro Gln Tyr Ala Asp 150 155 Cys Ala Ala Leu Pro Val Gly Ala Leu Ala Thr Glu Gln Trp Glu Glu

				165					170					175	
Asp	Pro	Ala	Val 180		Ala	Trp	Ser	Ile 185		Pro	Glu	Pro	Val 190		Gln
Glu	Glu	Ala 195	Ser	Ile	Trp	Pro	Phe 200	Glu	Gly	Leu	Gly	Gln 205	Leu	Gln	Pro
Pro	Ala 210	Val	Glu	Ile	Pro	Tyr 215	His	Glu	Ile	Leu	Trp 220	Arg	Glu	Trp	Glu
Asp 225	Phe	Ser	Thr	Gln	Pro 230	Asp	Ala	Gln	Gly	Leu 235	Lys	Ala	Gly	Asp	Gly 240
Pro	Gln	Phe	Gln	Phe 245	Thr	Leu	Met	Ser	Tyr 250	Asn	Ile	Leu	Ala	Gln 255	Asp
Leu	Met	Gln	Gln 260	Ser	Ser	Glu	Leu	Tyr 265	Leu	His	Cys	His	Pro 270	Asp	Ile
Leu	Asn	Trp 275	Asn	Tyr	Arg	Phe	Val 280	Asn	Leu	Met	Gln	Glu 285	Phe	Gln	His
Trp	Asp 290	Pro	Asp	Ile	Leu	Cys 295	Leu	Gln	Glu	Val	Gln 300	Glu	Asp	His	Tyr
Trp 305	Glu	Gln	Leu	Glu	Pro 310	Ser	Leu	Arg	Met	Met 315	Gly	Phe	Thr	Cys	Phe 320
Tyr	Lys	Arg	Arg	Thr 325	Gly	Cys	Lys	Thr	Asp 330	Gly	Cys	Ala	Val	Cys 335	Tyr
Lys	Pro	Thr	Arg 340	Phe	Arg	Leu	Leu	Cys 345	Ala	Ser	Pro	Val	Glu 350	Tyr	Phe
		355					360			Asn		365			
	370					375				Gly	380				
385		_			390					Tyr 395				_	400
				405					410	Leu				415	
			420					425		Pro			430		
_		435				_	440			Tyr		445		_	_
	450					455				Trp	460				
465	_				470		_		_	Lys 475					480
				485					490	Cys				495	
			500					505		Tyr			510		
Leu	Arg	Phe 515	Arg	Phe	Cys	Ser	Ile 520	Ala	Cys	Gln	Arg	Pro 525	Val	Gly	Leu
	530			_		535	_		_	Pro	540	-			
545					550					Ser 555					560
				565	_				570	Cys				575	
			580					585		Arg		•	590		
_		595					600			Tyr		605			
	610					615				His	620				
Gly 625	Thr	Leu	Lys	Leu	Leu 630	Gly	Arg	Leu	Ser	Leu 635	Leu	Ser	Glu	Glu	Ile 640

Leu Trp Ala Ala Asn Gly Leu Pro Asn Pro Phe Cys Ser Ser Asp His 645 650 655

Leu Cys Leu Leu Ala Ser Leu Gly Met Glu Val Thr Ala Pro * 660 670

<210> 1585 <211> 318 <212> PRT <213> Homo sapiens

<400> 1585 Met Met Cys Leu Lys Ile Leu Arg Ile Ser Leu Ala Ile Leu Ala Gly 10 Trp Ala Leu Cys Ser Ala Asn Ser Glu Leu Gly Trp Thr Arg Lys Lys 2.0 25 Ser Leu Val Glu Arg Glu His Leu Asn Gln Val Leu Leu Glu Gly Glu 40 Arg Cys Trp Leu Gly Ala Lys Val Arg Arg Pro Arg Ala Ser Pro Gln His His Leu Phe Gly Val Tyr Pro Ser Arg Ala Gly Asn Tyr Leu Arg 70 Pro Tyr Pro Val Gly Glu Glu Ile His His Thr Gly Arg Ser Lys Pro Asp Thr Glu Gly Asn Ala Val Ser Leu Val Pro Pro Asp Leu Thr 105 Glu Asn Pro Ala Gly Leu Arg Gly Ala Val Glu Glu Pro Ala Ala Pro 120 125 Trp Val Gly Asp Ser Pro Ile Gly Gln Ser Glu Leu Leu Gly Asp Asp 135 140 Asp Ala Tyr Leu Gly Asn Gln Arg Ser Lys Glu Ser Leu Gly Glu Ala 150 155 Gly Ile Gln Lys Gly Ser Ala Met Ala Ala Thr Thr Thr Ala Ile 165 170 Phe Thr Thr Leu Asn Glu Pro Lys Pro Glu Thr Gln Arg Arg Gly Trp 180 185 Ala Lys Ser Arg Gln Arg Gln Val Trp Lys Arg Arg Ala Glu Asp 200 205 Gly Gln Gly Asp Ser Gly Ile Ser Ser His Phe Gln Pro Trp Pro Lys 215 220 His Ser Leu Lys His Arg Val Lys Lys Ser Pro Pro Glu Glu Ser Asn 230 235 Gln Asn Gly Gly Glu Gly Ser Tyr Arg Glu Ala Glu Thr Phe Asn Ser 245 250 Gln Val Gly Leu Pro Ile Leu Tyr Phe Ser Gly Arg Arg Glu Arg Leu 265 Leu Leu Arg Pro Glu Val Leu Ala Glu Ile Pro Arg Glu Ala Phe Thr 280 285 Val Glu Ala Trp Val Lys Pro Glu Gly Gly Gln Asn Asn Pro Ala Ile 295 Ile Ala Gly Asn Thr Leu Leu Leu Gly Phe Leu Lys Ser * 310 315

<210> 1586 <211> 80

<212> PRT <213> Homo sapiens

<210> 1587 <211> 316 <212> PRT <213> Homo sapiens

<400> 1587 Met Phe Phe Gly Ser Ala Ala Leu Gly Thr Leu Thr Gly Leu Ile Ser 10 Ala Leu Val Leu Lys His Ile Asp Leu Arg Lys Thr Pro Ser Leu Glu 25 Phe Gly Met Met Ile Ile Phe Ala Tyr Leu Pro Tyr Gly Leu Ala Glu 40 Gly Ile Ser Leu Ser Gly Ile Met Ala Ile Leu Phe Ser Gly Ile Val 55 Met Ser His Tyr Thr His His Asn Leu Ser Pro Val Thr Gln Ile Leu 70 75 Met Gln Gln Thr Leu Arg Thr Val Ala Phe Leu Cys Glu Thr Cys Val 90 Phe Ala Phe Leu Gly Leu Ser Ile Phe Ser Phe Pro His Lys Phe Glu 105 110 Ile Ser Phe Val Ile Trp Cys Ile Val Leu Val Leu Phe Gly Arg Ala 120 125 Val Asn Ile Phe Pro Leu Ser Tyr Leu Leu Asn Phe Phe Arg Asp His 135 Lys Ile Thr Pro Lys Met Met Phe Ile Met Trp Phe Ser Gly Leu Arg 150 155 Gly Ala Ile Pro Tyr Ala Leu Ser Leu His Leu Asp Leu Glu Pro Met 170 Glu Lys Arg Gln Leu Ile Gly Thr Thr Thr Ile Val Ile Val Leu Phe 185 Thr Ile Leu Leu Gly Gly Ser Thr Met Pro Leu Ile Arg Leu Met 200 Asp Ile Glu Asp Ala Lys Ala His Arg Arg Asn Lys Lys Asp Val Asn · 220 215 Leu Ser Lys Thr Glu Lys Met Gly Asn Thr Val Glu Ser Glu His Leu 230 235 Ser Glu Leu Thr Glu Glu Glu Tyr Glu Ala His Tyr Ile Arg Arg Gln 245 250 Asp Leu Lys Gly Phe Val Trp Leu Asp Ala Lys Tyr Leu Asn Pro Phe

<210> 1588

<211> 53

<212> PRT

<213> Homo sapiens

<221> misc feature

<222> (1)...(53)

<223> Xaa = any amino acid or nothing

<400> 1588

 Met Cys Ser Leu Met Phe Gly Ser Ser Val Phe Val Cys Phe Pro Pro 1
 5
 10
 15

 Cys Val Pro Leu Pro Ala Pro His Ser Gly Gly Pro Pro His Arg Ala 20
 25
 30

 Gly Arg Ser Val Phe Ser Ala Met Lys Leu Gly Lys Xaa Arg Ser His 35
 40
 45

 Lys Glu Glu Pro Gln 50
 53

<210> 1589

<211> 437

<212> PRT

<213> Homo sapiens

<400> 1589

Met Leu Lys Val Ser Ala Val Leu Cys Val Cys Ala Ala Ala Trp Cys 10 Ser Gln Ser Leu Ala Ala Ala Ala Ala Val Ala Ala Gly Gly Arg 25 Ser Asp Gly Gly Asn Phe Leu Asp Asp Lys Gln Trp Leu Thr Thr Ile 40 Ser Gln Tyr Asp Lys Glu Val Gly Gln Trp Asn Lys Phe Arg Asp Glu 60 55 Val Glu Asp Asp Tyr Phe Arg Thr Trp Ser Pro Gly Lys Pro Phe Asp Gln Ala Leu Asp Pro Ala Lys Asp Pro Cys Leu Lys Met Lys Cys Ser 90 85 Arg His Lys Val Cys Ile Ala Gln Asp Ser Gln Thr Ala Val Cys Ile 105 Ser His Arg Arg Leu Thr His Arg Met Lys Glu Ala Gly Val Asp His 120 125 Arg Gln Trp Arg Gly Pro Ile Leu Ser Thr Cys Lys Gln Cys Pro Val 140 135 Val Tyr Pro Ser Pro Val Cys Gly Ser Asp Gly His Thr Tyr Ser Phe 155 150 Gln Cys Lys Leu Glu Tyr Gln Ala Cys Val Leu Gly Lys Gln Ile Ser

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170
             165
Val Lys Cys Glu Gly His Cys Pro Cys Pro Ser Asp Lys Pro Thr Ser
                    185
Thr Ser Arg Asn Val Lys Arg Ala Cys Ser Asp Leu Glu Phe Arg Glu
                       200
Val Ala Asn Arg Leu Arg Asp Trp Phe Lys Ala Leu His Glu Ser Gly
       215
                            220
Ser Gln Asn Lys Lys Thr Lys Thr Leu Leu Arg Pro Glu Arg Ser Arg
    230
                       235
Phe Asp Thr Ser Ile Leu Pro Ile Cys Lys Asp Ser Leu Gly Trp Met
       245 250
Phe Asn Arg Leu Asp Thr Asn Tyr Asp Leu Leu Asp Gln Ser Glu
   260 265
Leu Arg Ser Ile Tyr Leu Asp Lys Asn Glu Gln Cys Thr Lys Ala Phe
 275 280 285
Phe Asn Ser Cys Asp Thr Tyr Lys Asp Ser Leu Ile Ser Asn Asn Glu
                   295
Trp Cys Tyr Cys Phe Gln Arg Gln Gln Asp Pro Pro Cys Gln Thr Glu
                310
Leu Ser Asn Ile Gln Lys Arg Gln Gly Val Lys Lys Leu Leu Gly Gln
             325
                               330
Tyr Ile Pro Leu Cys Asp Glu Asp Gly Tyr Tyr Lys Pro Thr Gln Cys
                          345
His Gly Ser Val Gly Gln Cys Trp Cys Val Asp Arg Tyr Gly Asn Glu
                                         365
                       360
Val Met Gly Ser Arg Ile Asn Gly Val Ala Asp Cys Ala Ile Asp Phe
                    375
                                      380
Glu Ile Ser Gly Asp Phe Ala Ser Gly Asp Phe His Glu Trp Thr Asp
                390
                                  395
Asp Glu Asp Asp Glu Asp Asp Ile Met Asn Asp Glu Asp Glu Ile Glu
             405
                              410
Asp Asp Glu Asp Glu Gly Asp Asp Asp Gly Gly Asp Asp His
                          425
Asp Val Tyr Ile *
      435 436
```

<210> 1590

<211> 49

<212> PRT

<213> Homo sapiens

<400> 1590

 Met Phe Gln Ile Tyr Phe Ser Phe Cys Gln Leu Cys Phe Ile Trp Ser

 1
 5
 10
 15

 Cys Phe Phe Asn Ser Arg Glu Thr Phe Asn Glu Ile Tyr Lys Phe Phe
 20
 25
 30

 Leu Lys Ser Val Met Val Arg Lys Ile Phe Glu Cys His Lys Met Ser
 35
 40
 45
 48

<210> 1591

<211> 73

<212> PRT

<213> Homo sapiens

<400> 1591

<210> 1592

<211> 62

<212> PRT

<213> Homo sapiens

<400> 1592

 Met
 Tyr
 Phe
 Ser
 Leu
 Ile
 Phe
 Leu
 Val
 Phe
 Phe
 Phe
 Leu
 Pro
 15

 Leu
 Ser
 Ser
 Ser
 Ser
 Glu
 Pro
 Thr
 Ser
 Ser
 Ile
 Leu
 Gly
 Phe
 Ser

 Ser
 Ser
 Ser
 Ser
 Ser
 Ser
 Ser
 Phe
 Ser
 Phe
 Ser
 Phe
 Ser
 Phe
 Ser
 Leu
 Ser
 Ser
 Ala

 Ser
 Ser
 Ser
 Leu
 Ile
 Ser
 Phe
 Ser
 Phe
 Ser
 Phe
 Ser
 Lys
 *

 50
 Ser
 Ser
 Ser
 Ser
 Ser
 Phe
 Ser
 Phe
 Ser
 Phe
 Ser
 Lys
 *

<210> 1593

<211> 128

<212> PRT

<213> Homo sapiens

<400> 1593

Met Arg Ala Met Leu Gly Thr Cys Ala Leu Gly Gln Phe Phe Leu Ile 10 Met Gly Asn Thr Gln Arg Cys Asp Asp Phe Pro Thr Glu Ser Pro Pro 20 25 Ala Lys Thr Asn Val Ser Arg Ala Gly Leu Ser Pro Pro Cys Glu Ala 40 45 Leu His Gly Val Glu Ser Arg Gly Ser Cys Ser His Gly Lys Leu Gln 55 Ser Pro Pro Gly Arg Asp Trp Pro Gln Gly Asp Pro Gln Asp Arg Pro Lys Arg Arg Trp Gln Arg Pro Gly Pro Ala Gly Arg Gly Ala Pro Asp Pro Thr Pro Lys Gly Gln Gly Ala Ala Val Pro Pro Arg Ser Ala Ser 105 Met Phe Leu Ile His Lys Gln Met Trp Ala Tyr Gly Phe Gly Asp * 125

<210> 1594 <211> 46 <212> PRT <213> Homo sapiens

40

<210> 1595 <211> 86 <212> PRT <213> Homo sapiens

<210> 1596 <211> 69 <212> PRT <213> Homo sapiens

<210> 1597 <211> 56 <212> PRT <213> Homo sapiens

<400> 1597

<210> 1598 <211> 97 <212> PRT <213> Homo sapiens

<400> 1598 Met His Glu Ser Pro Leu Ala Trp Ala Ser Val His Leu Ser Ser Leu 10 Pro Leu Leu Cys Thr Ala Cys Ser Ser Pro Leu Met Gly Asn Ser Val 20 25 Leu Cys Arg Ala Pro Ala Asp Met Gly Leu Ala Trp Met Leu Leu Leu 40 Ser Glu Pro Arg Arg Val Val Pro Gly Ile Ala Ala Gln Val Leu Thr 55 60 Ala Leu Arg Arg Leu Leu Ser Gly Thr Leu Pro Ser Phe Pro Arg 70 75 Arg Lys Asn Pro Leu His Glu His Leu Leu Ala Phe Ile Val Arg Leu 90

<210> 1599 <211> 113 <212> PRT <213> Homo sapiens

65 70 75 80

Asp Pro Tyr His Leu Ser Arg Asp Leu Tyr Tyr Leu Thr Val Glu Ser 85 90 95

Ser Glu Lys Glu Ser Cys Arg Thr Pro Lys Val Val Asp Ile Pro Asp 100 105 110 112

<210> 1600 <211> 103 <212> PRT <213> Homo sapiens

<400> 1600 Met Gly Ala Trp Ala Trp Val Pro Thr Pro Ser Leu Cys Leu Cys His 1 5 10 Ser Thr Cys Leu Glu Phe Leu Leu Phe Leu Tyr Ile Leu Phe Tyr Cys 20 25 Ile Phe Glu Thr Val Ser Leu Ser Pro Arg Leu Glu Arg Ser Gly Ala 40 Ile Leu Ala Arg Cys Asn Leu Cys Leu Arg Gly Ser Ser Asp Ser Arg 55 Ala Leu Ala Ser Arg Val Ala Glu Thr Thr Gly Met His His His Ala 70 75 Trp Leu Ile Phe Ala Phe Leu Val Glu Thr Gly Phe His His Val Gly 85 90 Gln Ala Gly Leu Asn Ser * 100 102

<210> 1601 <211> 84 <212> PRT <213> Homo sapiens

<210> 1602 <211> 91 <212> PRT

<213> Homo sapiens

<400> 1602 Met Lys Thr Leu Pro Val Leu Val Leu Ser Leu Thr Leu Leu Thr Val 5 10 Phe Ser Glu Thr Ser Pro Ile Leu Thr Glu Lys Gln Ala Lys Gln Leu 20 25 Leu Arg Ser Arg Arg Gln Asp Arg Pro Ser Lys Pro Gly Phe Pro Asp 40 45 Glu Pro Met Arg Glu Tyr Met His His Leu Leu Ala Leu Glu His Arg 55 60 Ala Glu Glu Gln Phe Leu Glu His Trp Leu Asn Pro His Cys Lys Pro 70 75 His Cys Asp Arg Asn Arg Ile His Pro Val * 85

<210> 1603 <211> 69 <212> PRT

<213> Homo sapiens

Ile Leu Glu Val * 65 68

> <210> 1604 <211> 83 <212> PRT <213> Homo sapiens

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<210> 1605
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    <213> Homo sapiens
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     <223> Xaa = any amino acid or nothing
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Met Ser Thr Ile Ile Phe Gln Trp Pro Phe Met Leu Val Ser Leu His
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Arg Cys Arg Lys Leu Pro Arg Ala Leu Lys Asp Trp Gln Ala Phe Leu
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                                25
Asp Leu Lys Lys Ile Ile Asp Asp Phe Ser Glu Cys Cys Pro Leu Leu
       35
                            40
                                                45
Glu Tyr Met Gly Ser Lys Ala Met Met Glu Arg His Xaa Glu Arg Ile
                        5.5
Thr Thr Leu Thr Gly His Ser Leu Asp Val Gly Asn Glu Ser Phe Lys
                                        75
Leu Arg Asn Ile Met Glu Ala Pro Leu Leu Xaa Tyr Lys Glu Glu Ile
Glu Val Glu Tyr Asp Val Met Glu Asp Cys Lys Val Ser Trp
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   <210> 1606
    <211> 72
    <212> PRT
    <213> Homo sapiens
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<400> 1606

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 Gly
 Thr
 Val
 Thr
 Met
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 Leu
 Trp
 His
 Ala
 Ser
 Asn
 Trp

 Asp
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 Gln
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 Pro
 Leu
 Val
 Glu
 Leu
 Thr
 Pro
 Val
 Arg

 Asp
 Val
 Ser
 Gly
 Leu
 Thr
 Ala
 Phe
 Leu
 Ala
 Arg
 Asp
 Met
 Asn

 Leu
 Leu
 Ser
 Gly
 Asn
 Val
 Asn
 Thr
 Met
 Asn
 Gly
 Glu
 Ser
 Ile
 Ile
 Ala

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 55
 55
 60
 60

 Ile
 Thr
 Met
 Leu
 Ala
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 65
 70
 71
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<210> 1607 <211> 59 <212> PRT <213> Homo sapiens

 Phe
 Leu
 Leu
 Ser
 Phe
 Ile
 Ser
 Tyr
 Phe
 Cys
 Leu
 Pho
 Pro
 Cys
 Ser

 Asn
 Leu
 Pro
 Lys
 Val
 Ile
 Ala
 Ile
 Phe
 Asn
 Ile
 Val
 Leu
 Ile
 Leu
 Ile
 Leu
 Ser

 Ile
 Val
 Phe
 Arg
 Glu
 Ile
 Thr
 Asp
 Thr
 Tyr
 *

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 55
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 58

<210> 1608 <211> 118 <212> PRT <213> Homo sapiens

<400> 1608 Met Leu Val Thr Asp Thr Glu Ala Phe Trp Gln Pro Gln Pro Trp Phe 10 Val Val Leu Thr Ala Thr Gly Ala Leu Leu Leu Ala Leu Gly 25 Trp Leu Leu Gly Arg Leu Leu Gln Gly Leu Ala Gln Leu Leu Gln Ala 40 Pro Ser Lys Pro Ala Gln Ala Leu Leu Leu Asn Ser Ile Gln Gly Thr Glu Gly Ser Ile Glu Gly Phe Leu Glu Ala Pro Lys Met Glu Met Ser 75 Gln Ala Pro Ser Ser Val Met Ser Leu Gln His Phe Asp Gly Arg Thr 90 85 Gln Asp Ser Arg Thr Gly Arg Asp Tyr Leu Val Asn Thr His Thr Gly 105 100 Ala Arg Arg Trp Leu * 115 117

<210> 1609 <211> 50 <212> PRT <213> Homo sapiens

<400> 1609
Met Val Ile Gly Ser Leu His Thr Phe Thr Leu Leu Ala Ala Ser Ser

Leu Val Asp Thr Pro Lys Gln Ile Gln Leu Leu Met Gln Asn Leu Met

20

25

Asn Asp Pro Arg Lys Glu Val Lys Ile Leu Ala Ile Gln Asp Leu Lys

Ash Asp Pro Arg Lys Giu Vai Lys lie Leu Ala lie Gin Asp Leu I 35 40 45 Leu Leu

Leu Leu 50

> <210> 1610 <211> 50 <212> PRT <213> Homo sapiens

<210> 1.611 <211> 56 <212> PRT

<213> Homo sapiens

<400> 1611

 Met
 Ser
 Phe
 Gln
 Ala
 Phe
 Val
 Phe
 Leu
 Met
 Ile
 Gly
 Trp
 Leu
 His
 Pro

 Asp
 Pro
 Arg
 Leu
 Met
 Thr
 Gln
 Arg
 Ser
 Cys
 Gly
 Pro
 His
 Pro
 Glu
 Val

 20
 20
 25
 30
 30
 Asn
 Asn
 Asn
 His
 Pro
 Ser
 His
 Pro
 Tyr
 Asp
 Ile
 Pro
 Asn

 Gln
 Ser
 Ala
 Pro
 Pro
 Leu
 Pro
 *
 45

<210> 1612 <211> 75 <212> PRT <213> Homo sapiens

<210> 1613 <211> 192 <212> PRT <213> Homo sapiens

Arg Lys Ser Asp Pro Lys Arg Phe Gln Asn Ile Phe Thr Thr Ile Phe Thr Leu Phe Thr Leu Leu Thr Leu Asp Asp Trp Ser Leu Ile Tyr Met Asp Ser Arg Ala Gln Gly Ala Trp Tyr Ile Ile Pro Ile Leu Ile Ile 55 Tyr Ile Ile Ile Gln Tyr Phe Ile Phe Leu Asn Leu Val Ile Thr Val 75 70 Leu Val Asp Ser Phe Gln Thr Ala Leu Phe Lys Gly Leu Glu Lys Ala 90 Lys Gln Glu Arg Ala Ala Arg Ile Gln Glu Lys Leu Leu Glu Asp Ser 105 Leu Thr Glu Leu Arg Ala Ala Glu Pro Lys Glu Val Ala Ser Glu Gly 125 120 Thr Met Leu Lys Arg Leu Ile Glu Lys Lys Phe Gly Thr Met Thr Glu 135 140 Lys Gln Gln Glu Leu Leu Phe His Tyr Leu Gln Leu Val Ala Ser Val 155 - 160 150 Glu Gln Glu Gln Gln Lys Phe Arg Ser Gln Ala Ala Val Ile Asp Glu 170 165 Ile Val Asp Thr Thr Phe Glu Ala Gly Glu Glu Asp Phe Arg Asn * 185 180

<210> 1614 <211> 153 <212> PRT <213> Homo sapiens

<400> 1614 Met Asp Leu Val Gln Phe Phe Val Thr Phe Phe Ser Cys Phe Leu Ser Leu Leu Leu Val Ala Ala Val Val Trp Lys Ile Lys Gln Thr Cys Trp 25 Ala Ser Arg Arg Arg Glu Gln Leu Leu Arg Glu Arg Gln Gln Met Ala 40 Ser Arg Pro Phe Ala Ser Val Asp Val Ala Leu Glu Val Gly Ala Glu 55 Gln Thr Glu Phe Leu Arg Gly Pro Leu Glu Gly Ala Pro Lys Pro Ile 75 70 Ala Ile Glu Pro Cys Ala Gly Asn Arg Ala Ala Val Leu Thr Val Phe 90 85 Leu Cys Leu Pro Arg Gly Ser Ser Gly Ala Pro Pro Pro Gly Gln Ser 105 Gly Leu Ala Ile Ala Ser Ala Leu Ile Asp Ile Ser Gln Gln Lys Ala 125 120 Ser Asp Ser Lys Asp Lys Thr Ser Gly Val Arg Asn Arg Lys His Leu 140 135 Ser Thr Arg Gln Gly Thr Cys Val * 150 152 145

<210> 1615 <211> 135 <212> PRT <213> Homo sapiens

<400> 1615 Met His Trp Leu Arg Ala Ser Ala Gly Ser Leu Leu Met Val Pro Leu 10 Met Thr Asp Leu His Glu Leu Ala Leu Pro Pro Ala Ser Leu Arg Thr 25 20 Val Val Lys Glu Asn Met Cys Val Leu Pro Phe Pro Val Lys Thr Ser 40 45 Gly Arg Ser Leu Thr Gly Ser Ala Trp Ser Arg Phe His Leu Pro Cys 55 His Leu Arg Pro Gly Asp Arg Leu Pro Cys His Cys Leu Gly Lys Phe 70 Arg Lys Arg Val Ala Lys Trp Cys Ile Arg Lys Asn Met Ala Arg Ser 85 90 Pro His Leu Leu Gly Gly Arg Pro Asn Ser Thr Ser Gly Pro Leu Cys 105 Asp Phe Pro Ala Pro Ser Lys Gln Val Thr Pro Leu Leu Trp Val Ser 120 Val Ser Leu Pro Ile Lys * 130 134

<210> 1616 <211> 60 <212> PRT

<213> Homo sapiens

<210> 1617 <211> 49 <212> PRT <213> Homo sapiens

<210> 1618 <211> 95 <212> PRT <213> Homo sapiens

<400> 1618 Met Trp Thr Val Leu Trp His Arg Phe Ser Met Val Leu Arg Leu Pro 5 10 Glu Glu Ala Ser Ala Gln Glu Gly Glu Leu Ser Leu Ser Pro Pro 20 25 Ser Pro Glu Pro Asp Trp Thr Leu Ile Ser Pro Gln Gly Met Ala Ala 40 Leu Leu Ser Leu Ala Met Ala Thr Phe Thr Gln Glu Pro Gln Leu Cys 55 Leu Ser Cys Leu Ser Gln His Gly Ser Ile Leu Met Ser Ile Leu Lys 70 75 His Leu Cys Pro Ser Phe Leu Asn Gln Leu Arg Gln Ala * 85 90

<210> 1619 <211> 54 <212> PRT <213> Homo sapiens

<210> 1620 <211> 71 <212> PRT <213> Homo sapiens

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<210> 1621
     <211> 90
     <212> PRT
     <213> Homo sapiens
    <221> misc_feature
     <222> (1)...(90)
     <223> Xaa = any amino acid or nothing
    <400> 1621
Met Asp His Lys Ser Leu Trp Ala Gly Val Glu Val Leu Leu Leu Leu
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Gln Gly Gly Ser Ala Tyr Lys Leu Val Cys Tyr Phe Thr Asn Trp Ser
                                25
Gln Asp Arg Gln Glu Pro Gly Lys Phe Thr Pro Glu Asn Ile Asp Pro
                            40
Phe Leu Cys Ser His Leu Ile Tyr Ser Phe Ala Ser Ile Glu Asn Asn
                        55
Lys Val Ile Ile Arg Thr Pro Xaa Phe Phe Pro Leu Pro Leu Gly His
                    70 ·
Arg Leu Gln Thr Ile Asn Pro Arg Leu *
                85
     <210> 1622
     <211> 53
     <212> PRT
     <213> Homo sapiens
    <400> 1622
Met Gln Cys Ala Ile Cys Ile Leu Leu Tyr Leu Leu Asn Lys Lys Thr
                                    10
Val Trp Arg Cys Ser Arg Ile His His Asn Asn Thr Val Val Leu Thr
                                25
Arg Glu Ser Ser Pro Phe Leu Thr Thr Cys Thr Leu Ser Ser Val Leu
       35
                           40
Leu Thr Lys Ala *
    50 52
    <210> 1623
    <211> 978
    <212> PRT
    <213> Homo sapiens
    <400> 1623
Met Pro Ala Arg Arg Leu Leu Leu Leu Leu Thr Leu Leu Pro Gly
                                   10
Leu Gly Ile Phe Gly Ser Thr Ser Thr Val Thr Leu Pro Glu Thr Leu
                                25
Leu Phe Val Ser Thr Leu Asp Gly Ser Leu His Ala Val Ser Lys Arg
```

Thr	Gly	Ser	Ile	Lys	Trp	Thr	Leu	Lys	Glu	Asp	Pro	Val	Leu	Gln	Val
D	50	TT -	777	a 1	G3	55	27-	To 1	•	5	60	D	7	70	al
65					70					75		Pro		*	80
Ser	Leu	Tyr	Thr	Leu 85	Gly	Ser	Lys	Asn	Asn 90	Glu	Gly	Leu	Thr	Lys 95	Leu
Pro	Phe	Thr	Ile 100	Pro	Glu	Leu	Val	Gln 105	Ala	Ser	Pro	Cys	Arg 110	Ser	Ser
Asp	Gly	Ile 115	Leu	Tyr	Met	Gly	Lys 120	Lys	Gln	Asp	Ile	Trp 125	Tyr	Val	Ile
Asp	Leu 130		Thr	Gly	Glu	Lys 135		Gln	Thr	Leu	Ser 140	Ser	Ala	Phe	Ala
Asp 145	-	Leu	Cys	Pro	Ser 150		Ser	Leu	Leu	Tyr 155		Gly	Arg	Thr	Glu 160
-	Thr	Ile	Thr	Met 165		Asp	Thr	Lys	Thr 170		Glu	Leu	Arg	Trp 175	
Ala	Thr	Tyr	Phe 180		Tyr	Ala	Ala	Ser 185		Pro	Glu	Asp	Asp		Asp
Tyr	Lys	Met 195		His	Phe	Val	Ser 200		Gly	Asp	Gly	Leu 205		Val	Thr
Val	Asp 210		Glu	Ser	Gly	Asp 215		Leu	Trp	Ile	Gln 220	Asn	Tyr	Ala	Ser
Pro 225		Val	Ala	Phe	Tyr 230		Trp	Gln	Arg			Leu	Arg	Lys	
	His	Ile	Asn			Val	Glu	Thr		235 Arg	Tyr	Leu	Thr		240 Met
Ser	Gly	Glu		245 Gly	Arg	Ile	Thr	_	250 Trp	Lys	Tyr	Pro		255 Pro	Lys
Glu	Thr		260 Ala	Lys	Ser	Lys		265 Thr	Pro	Thr	Leu	Tyr	270 Val	Gly	Lys
Tyr		275 Thr	Ser	Leu	Tyr		280 Ser	Pro	Ser	Met		285 His	Glu	Gly	Val
	290 Val	Val	Pro	Arg	_	295 Ser	Thr	Leu	Pro		300 Leu	Glu	Gly	Pro	
305 Thr	Asp	Glv	Val	Thr	310 Tle	Glv	Asn	Lave	Glv	315	Cve	Val	Tle	Thr	320 Pro
				325			_	_	330		_			335	
Ser	Thr	Asp	Val 340	Lys	Phe	Asp	Pro	Gly 345	Leu	Lys	Ser	Lys	Asn 350	Lys	Leu
Asn	Tyr	Leu 355	Arg	Asn	Tyr	Trp	Leu 360	Leu	Ile	Gly	His	His 365	Glu	Thr	Pro
Leu	Ser 370	Ala	Ser	Thr	_	Met 375	Leu	Glu	Arg	Phe	Pro 380	Asn	Asn	Leu	Pro
Lys 385	His	Arg	Glu	Asn	Val 390	Ile	Pro	Ala	Asp	Ser 395	Glu	Lys	Lys	Ser	Phe 400
Glu	Glu	Val	Ile	Asn 405	Leu	Val	Asp	Gln	Thr 410	Ser	Glu	Asn	Ala	Pro 415	Thr
Thr	Val	Ser	Arg 420	Asp	Val	Glu	Glu	Lys 425	Pro	Ala	His	Ala	Pro 430	Ala	Arg
Pro	Glu	Ala 435	Pro	Val	Asp	Ser	Met 440	Leu	Lys	Asp	Met	Ala 445	Thr	Ile	Ile
Leu	Ser 450	Thr	Phe	Leu	Leu	Ile 455	Gly	Trp	Val	Ala	Phe 460	Ile	Ile	Thr	Tyr
Pro 465		Ser	Met	His	Gln 470		Gln	Gln	Leu	Gln 475		Gĺn	Gln	Phe	Gln 480
	Glu	Leu	Glu	Lys 485		Gln	Leu	Leu	Gln 490		Gln	Gln	Gln	Gln 495	
Pro	Phe	His	Pro 500		Gly	Asp	Thr	Ala 505		Asp	Gly	Glu	Leu 510		Asp
Thr	Ser	Gly		Tyr	Ser	Glu	Ser		Gly	Thr	Ser	Ser		Ser	Thr

		515					520					525			
Ser	Pro 530		Ala	Ser	Asn	His 535		Leu	Cys	Ser	Gly 540		Ser	Ala	Ser
Lys 545	Ala	Gly	Ser	Ser	Pro 550		Leu	Glu	Gln	Asp 555		Gly	Asp	Glu	Glu 560
Thr	Ser	Val	Val	Ile 565	Val	Gly	Lys	Ile	Ser 570	Phe	Cys	Pro	Lys	Asp 575	Val
			Gly 580					585		_			590		
		595	Val				600					605			
	610		Glu			615					620				
625			Tyr		630			_		635					640
			Leu	645					650		_			655	_
			His 660		_			665					670		
		675	Leu				680					685			
	690		His			695					700			_	_
705			Met		710					715					720
			Ser Pro	725					730					735	
			740 Asp					745					750		
		755	Ser				760					765			
	770		Gly			775	_	_			780				
785					790					795					800
			Ile	805					810					815	
			Arg 820					825					830		
		835	Glu				840					845			
	850		Glu			855					860				
865			Ala		870					875					880
			Thr	885					890					895	
			100 Leu					905		-	_		910	_	
		915	Ala				920			_		925			
	930		Tyr			935					940	•			
1yr 945	arg	Ala	Met	GIu	Leu 950	Cys	Ser	His	Glu	Arg 955	Leu	Phe	GIN	Pro	Tyr 960
Tyr	Phe	His	Glu	Pro 965		Glu	Pro	Gln	Pro 970		Val	Thr	Pro	Asp 975	
Leu 977	*														

<210> 1624 <211> 56 <212> PRT <213> Homo sapiens

<210> 1625 <211> 146 <212> PRT <213> Homo sapiens

<400> 1625 Met Glu Leu Ala Leu Leu Cys Gly Leu Val Val Met Ala Gly Val Ile 10 Pro Ile Gln Gly Gly Ile Leu Asn Leu Asn Lys Met Val Lys Gln Val 25 Thr Gly Lys Met Pro Ile Leu Ser Tyr Trp Pro Tyr Gly Cys His Cys 40 Gly Leu Gly Gly Arg Gly Gln Pro Lys Asp Ala Thr Asp Trp Cys Cys 55 Gln Thr His Asp Cys Cys Tyr Asp His Leu Lys Thr Gln Gly Cys Gly 70 Ile Tyr Lys Asp Tyr Tyr Arg Tyr Asn Phe Ser Gln Gly Asn Ile His 90 Cys Ser Asp Lys Gly Ser Trp Cys Glu Gln Gln Leu Cys Ala Cys Asp 105 Lys Glu Val Ala Phe Cys Leu Lys Arg Asn Leu Asp Thr Tyr Gln Lys 120 125 Arg Leu Arg Phe Tyr Trp Arg Pro His Cys Arg Gly Gln Thr Pro Gly 130 135 Cys * 145

<210> 1626 <211> 385 <212> PRT <213> Homo sapiens

<400> 1626
Met Glu Phe Gly Leu Ser Trp Leu Phe Leu Val Ala Ile Leu Lys Gly

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Val Gln Cys Glu Val Gln Leu Val Glu Ser Gly Gly Leu Val Gln
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Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe
                           40
Ser Ser Tyr Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu
                       55
                                           60
Glu Trp Val Ser Gly Ile Gly Gly Ser Gly Ser Ser Thr Tyr Tyr Ala
                    70
                                        75
Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Gln Asn
                                    90
Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val
                               105
Tyr Tyr Cys Ala Lys Ser His Pro Ala Tyr Tyr Tyr Gly Ser Gly Ser
                          120
                                              125
Tyr Ser Ser His Tyr Tyr Tyr Tyr Gly Met Asp Val Trp Gly Gln
                       135
                                           140
Gly Thr Thr Val Thr Val Ser Ser Gly Asp Gly Ser Ser Gly Gly Ser
                   150
                                       155
Gly Gly Ala Ser Thr Gly Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
                                   170
Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
                               185
Gln Ser Val Ser Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
                           200
Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Ser Ser Arg Ala Thr Gly
                       215
Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu
                   230
                                       235
Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
               245
                                   250
Gln Tyr Gly Ser Ser Pro Thr Thr Phe Gly Gln Gly Thr Lys Val Glu
                               265
                                                   270
Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
                          280
Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
                       295
Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
                  310
                                       315
Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
                                  330
              325
Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
                              345
           340
Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Ser Gly Ala
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Leu Ser Phe Ala Arg Ser Gln Arg Ser Phe Gln Pro Gly Glu Ser Val
                                           380
                       375
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<210> 1627 <211> 101

<212> PRT

<213> Homo sapiens

<400> 1627

 Met
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 His
 Cys
 Thr
 Ile
 Ile
 Pro
 Leu
 Ser
 Phe
 Cys
 Val
 His
 Arg

 Leu
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 Ala
 Pro
 Leu
 Asp
 Ala
 Tyr
 Phe
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<210> 1628 <211> 71 <212> PRT <213> Homo sapiens

<210> 1629 <211> 112 <212> PRT <213> Homo sapiens

<400> 1629

Met Ala His Tyr Lys Thr Glu Gln Asp Asp Trp Leu Ile Ile Tyr Leu 10 Lys Tyr Leu Leu Phe Val Phe Asn Phe Phe Phe Trp Val Gly Gly Ala 20 25 Ala Val Leu Ala Val Gly Ile Trp Thr Leu Val Glu Lys Ser Gly Tyr 40 Leu Ser Val Leu Ala Ser Ser Thr Phe Ala Ala Ser Ala Tyr Ile Leu 55 Ile Phe Ala Gly Val Leu Val Met Val Thr Gly Phe Leu Gly Phe Gly 75 Ala Ile Leu Trp Glu Arg Lys Gly Cys Leu Ser Thr Tyr Phe Cys Leu 85 90 Leu Leu Val Ile Phe Leu Asp Glu Leu Glu Ala Gly Val Leu Ala His 100

10

<210> 1630 <211> 47 <212> PRT <213> Homo sapiens <400> 1630 Met Trp Pro Gln Leu Lys Ser Phe Phe Leu Ile Pro Thr Gln Ile 1 5

20 25 Arg Phe Val Trp Val Ser Met Pro Glu Leu Ile Gly Ala Ser 40

His Phe Asn Leu Thr Asn Leu Pro Ser Trp Arg Arg Glu Leu Arg

<210> 1631 <211> 79 <212> PRT <213> Homo sapiens

<400> 1631 Met Tyr Met Trp Ser Gly Leu Leu Gly Ser Lys Trp Thr Leu Val Tyr 1 5 10 15 Ser His Phe Leu Asn Met Ala Pro Ala Ser Phe Ser His Tyr Gln Ala 25 Ser Leu Pro Leu Leu Glu His Asp Thr Leu Ser Ser Arg Val His 40 Ser Tyr Gln Cys Pro Gly Phe Phe Cys Phe Phe Pro Ser Val Leu Glu 55 60 Phe Ser Gln Leu Gln Lys Thr Tyr Ser Leu Cys Leu Pro Phe * 75

<210> 1632 <211> 48 <212> PRT <213> Homo sapiens

<400> 1632 Met Phe Met Cys Arg Leu Leu Trp Ala Thr Gly Ala Tyr Gly Phe 5 10 Leu Gly Asp Asp Val Glu Tyr Thr Ser Val Leu Pro His Gln Lys Gly 25 Lys Glu Ala Trp Val Phe Ile Cys Gln Leu Pro Phe Ile Ile Gly * 40

<210> 1633 <211> 58 <212> PRT

<213> Homo sapiens

<210> 1634 <211> 55 <212> PRT <213> Homo sapiens

<210> 1635 <211> 78 <212> PRT <213> Homo sapiens

<210> 1636 <211> 51 <212> PRT <213> Homo sapiens

<210> 1637 <211> 123 <212> PRT <213> Homo sapiens

---- crome capitons

<400> 1637 Met Gln Gln Met Met Trp Ala Gly Leu Leu Cys Pro Gln Leu Glu Trp 10 Leu Gln Gly Arg Ala Cys Arg Pro Cys Gly Leu Leu Ala Ser Asp Ala 25 Ala Ala Leu Trp Phe Arg Gly Gly Ile Ser Ala Trp Glu Asp Ser Cys 40 Ala Val Ser Asn Ile Arg His Glu Ala Tyr Asn Cys His Leu Ser Val 55 60 Phe Leu Asn Arg Cys Ala Asn Glu Leu Thr Val Gln Phe Leu Ile Ile 70 75 Leu Ala Phe Gln Ile Met Leu Ser Cys Ala Val Ile Ala Pro Ala Val 85 90 Pro Val Phe Gln Arg Leu Thr Leu Lys Arg Ser Gly Arg Thr Ser Leu 100 105 Gly Ser Thr Gly Arg Leu His Phe Cys Lys * 120

<210> 1638 <211> 69 <212> PRT <213> Homo sapiens

<210> 1639

<211> 92 <212> PRT <213> Homo sapiens

<210> 1640 <211> 58 <212> PRT <213> Homo sapiens

<210> 1641 <211> 459 <212> PRT <213> Homo sapiens

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Arg Ile Val Gln Leu Ile Gln Asp Thr Arg Ile His Ile Leu Pro Ser
                    120
Met Asn Pro Asp Gly Tyr Glu Val Ala Ala Ala Gln Gly Pro Asn Lys
                      135
                                         140
Pro Gly Tyr Leu Val Gly Arg Asn Ala Asn Gly Val Asp Leu Asn
                  150
                                      155
Arg Asn Phe Pro Asp Leu Asn Thr Tyr Ile Tyr Tyr Asn Glu Lys Tyr
                       170
              165
Gly Gly Pro Asn His His Leu Pro Leu Pro Asp Asn Trp Lys Ser Gln
                   185
          180
Val Glu Pro Glu Thr Arg Ala Val Ile Arg Trp Met His Ser Phe Asn
                         200
                                . 205
Phe Val Leu Ser Ala Asn Leu His Gly Gly Ala Val Val Ala Asn Tyr
                      215
                                         220
Pro Tyr Asp Lys Ser Phe Glu His Arg Val Arg Gly Val Arg Arg Thr
                  230
                                      235
Ala Ser Thr Pro Thr Pro Asp Asp Lys Leu Phe Gln Lys Leu Ala Lys
                                  250
Val Tyr Ser Tyr Ala His Gly Trp Met Phe Gln Gly Trp Asn Cys Gly
           260
                              265
Asp Tyr Phe Pro Asp Gly Ile Thr Asn Gly Ala Ser Trp Tyr Ser Leu
                          280
Ser Lys Gly Met Gln Asp Phe Asn Tyr Leu His Thr Asn Cys Phe Glu
                      295
                                          300
Ile Thr Leu Glu Leu Ser Cys Asp Lys Phe Pro Pro Glu Glu Glu Leu
                   310
                                      315
Gln Arg Glu Trp Leu Gly Asn Arg Glu Ala Leu Ile Gln Phe Leu Glu
               325
                                  330
Gln Val His Gln Gly Ile Lys Gly Met Val Leu Asp Glu Asn Tyr Asn
           340
                              345
Asn Leu Ala Asn Ala Val Ile Ser Val Ser Gly Ile Asn His Asp Val
                           360
Thr Ser Gly Asp His Gly Asp Tyr Phe Arg Leu Leu Pro Gly Ile
                       375
                                          380
Tyr Thr Val Ser Ala Thr Ala Pro Gly Tyr Asp Pro Glu Thr Val Thr
                   390
                                      395
Val Thr Val Gly Pro Ala Glu Pro Thr Leu Val Asn Phe His Leu Lys
               405
                                  410
Arg Ser Ile Pro Gln Val Ser Pro Val Arg Arg Ala Pro Ser Arg Arg
                              425
His Gly Val Arg Ala Lys Val Gln Pro Gln Pro Arg Lys Lys Glu Met
                          440
Glu Met Arg Gln Leu Gln Arg Gly Pro Ala
                       455
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<210> 1642

<211> 144

<212> PRT

<213> Homo sapiens

<400> 1642

Met Ala Arg Cys Thr Leu Thr Leu Leu Lys Thr Met Leu Thr Glu Leu 1 5 10 15

Leu Arg Gly Gly Ser Phe Glu Phe Lys Asp Met Arg Val Pro Ser Ala 20 25 30

Leu Val Thr Leu His Met Leu Leu Cys Ser Ile Pro Leu Ser Gly Arg 40 Leu Asp Ser Asp Glu Gln Lys Ile Gln Asn Asp Ile Ile Asp Ile Leu Leu Thr Phe Thr Gln Gly Val Asn Glu Lys Leu Thr Ile Ser Glu Glu 70 75 Thr Leu Ala Asn Asn Thr Trp Ser Leu Met Leu Lys Glu Val Leu Ser 85 90 Ser Ile Leu Lys Val Pro Glu Gly Phe Phe Ser Gly Leu Ile Leu Leu 105 Ser Glu Leu Pro Leu Pro Leu Pro Met Gln Thr Thr Gln Val Ser 120 Leu Pro Tyr Asn Met His Leu Ile Asn Asp Cys Ser Asn Thr Phe * 135 140

<210> 1643 <211> 70

<212> PRT

<213> Homo sapiens

<400> 1643

<210> 1644 <211> 82

<212> PRT

<213> Homo sapiens

<400> 1644

 Met Gly Met Gly Thr
 Leu Ile Ile Met Asn Val Trp Val Leu Phe Ile

 1
 5

 Pro Thr Arg Leu Arg Ile Asp Gln Gln Pro Val His Ile Lys Pro Ser

 20
 25

 30

 Met Arg Val Leu Asp Lys Trp Val Ser Ala Phe Val His Lys Gly Phe

 35
 40

 45

 Thr Trp Gly Thr Ser Glu Arg Ile Asn Thr Gly Ser Ser Ser Asp Ile

 50
 55

 60

 Thr Leu Gly Ile Leu Asn Lys Cys Gly Trp Ala Val Phe Cys Ala Ala

 65
 70

 75
 80

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<210> 1645
<211> 256
<212> PRT
<213> Homo sapiens
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<400> 1645 Met Ala Ala Leu Thr Val Thr Leu Met Val Leu Ser Ser Pro Leu Ala 1 5 10 Leu Ala Gly Asp Thr Gln Pro Arg Phe Leu Trp Gln Gly Lys Tyr Lys 25 Cys His Phe Phe Asn Gly Thr Glu Arg Val Gln Phe Leu Glu Arg Leu 40 Phe Tyr Asn Glu Glu Phe Val Arg Phe Asp Ser Asp Val Gly Glu 55 Tyr Arg Ala Val Thr Glu Leu Gly Arg Pro Val Ala Glu Ser Trp Asn 70 75 Ser Gln Lys Asp Ile Leu Glu Asp Arg Gly Gln Val Asp Thr Val 90 85 Cys Arg His Asn Tyr Gly Val Gly Glu Ser Phe Thr Val Gln Arg Arg 105 Val His Pro Glu Val Thr Val Tyr Pro Ala Lys Thr Gln Pro Leu Gln 120 125 His His Asn Leu Leu Val Cys Ser Val Ser Gly Phe Tyr Pro Gly Ser 135 Ile Glu Val Arg Trp Phe Arg Asn Gly Gln Glu Glu Lys Ala Gly Val 150 155 Val Ser Thr Gly Leu Ile Gln Asn Gly Asp Trp Thr Phe Gln Thr Leu 170 Val Met Leu Glu Thr Val Pro Arg Ser Gly Glu Val Tyr Thr Cys Gln 185 Val Glu His Pro Ser Val Met Ser Pro Leu Thr Val Glu Trp Arg Ala 200 Arg Ser Glu Ser Ala Gln Ser Lys Met Leu Ser Gly Val Gly Gly Phe 215 Val Leu Gly Leu Leu Phe Leu Gly Ala Gly Leu Phe Ile Tyr Phe Arg 230 235 Asn Gln Lys Gly His Ser Gly Leu Gln Pro Thr Gly Phe Leu Ser *

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<210> 1646
<211> 263
<212> PRT
<213> Homo sapiens
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Asp Asp Gly Arg Arg Lys Pro Gly Ile Gly Gly Arg Glu Arg Trp Asn 85 90 His Val Thr Thr Thr Lys Arg Pro Val Thr Thr Arg Ala Pro Ala 105 Asn Thr Leu Gly Asn Asp Phe Asp Leu Ala Asp Ala Leu Asp Asp Arg 120 Asn Asp Arg Asp Gly Arg Arg Lys Pro Ile Ala Gly Gly Gly 135 Phe Ser Asp Lys Asp Leu Glu Asp Ile Val Gly Gly Glu Tyr Lys 150 155 Pro Asp Lys Gly Lys Gly Asp Gly Arg Tyr Gly Ser Asn Asp Asp Pro 165 170 Gly Ser Gly Met Val Ala Glu Pro Gly Thr Ile Ala Gly Val Ala Ser 180 185 Ala Leu Ala Met Ala Leu Ile Gly Ala Val Ser Ser Tyr Ile Ser Tyr 200 205 Gln Gln Lys Lys Phe Cys Phe Ser Ile Gln Gln Gly Leu Asn Ala Asp 215 220 Tyr Val Lys Gly Glu Asn Leu Glu Ala Val Val Cys Glu Glu Pro Gln 230 235 Val Lys Tyr Ser Thr Leu His Thr Gln Ser Ala Glu Pro Pro Pro 250 Pro Glu Pro Ala Arg Ile * 260 262

<210> 1647

<211> 74

<212> PRT

<213> Homo sapiens

<400> 1647

 Met
 Tyr
 Leu
 Cys
 Trp
 Leu
 Tyr
 Ile
 Met
 Gly
 Val
 Leu
 Gly
 Ala
 Ser

 Cys
 Asn
 Trp
 His
 Val
 Gly
 Val
 Pro
 Phe
 Pro
 Gly
 Thr
 His
 Trp
 Pro
 Arg

 Ser
 Gln
 Asn
 His
 Leu
 Leu
 Trp
 Val
 Tyr
 Asn
 His
 Leu
 Asn
 Glu
 Leu
 Pro

 Val
 Pro
 Ala
 Gly
 Arg
 Ser
 Ser
 Glu
 Gln
 Leu
 Tyr
 Leu
 Arg
 Tyr
 Thr
 Gly
 Tyr
 Thr
 Glu

 Lys
 Tyr
 Gly
 Arg
 Arg
 Gly
 Arg
 Lys
 Ala
 *

 65
 Tyr
 Arg
 Arg
 Arg
 Lys
 Ala
 *

<210> 1648

<211> 58

<212> PRT

<213> Homo sapiens

<400> 1648

Met Gly Leu Cys Gly Met Trp Val Leu Thr Ala Phe Leu Cys Glu Pro 1 $$ 5 $$ 10 $$ 15 Met Gly Phe Arg His Arg Val Cys Pro His Arg Cys Val Arg Gly Ser 20 $$ 25 $$ 30 Gly Arg Gly Ser Gly Cys Glu Cys Val Thr Met Trp Pro Cys Gly Ile

35 40 45 Asn Ala Met Thr Gly Gly Phe Trp Val * 50 55 57

<210> 1649 <211> 90 <212> PRT <213> Homo sapiens

<210> 1650 <211> 113 <212> PRT <213> Homo sapiens

<400> 1650 Met Ala Leu Gly Val Pro Ile Ser Val Tyr Leu Leu Phe Asn Ala Met 1 5 10 15 Thr Ala Leu Thr Glu Glu Ala Ala Val Thr Val Thr Pro Pro Ile Thr Ala Gln Gln Gly Asn Trp Thr Val Asn Lys Thr Glu Ala Asp Asn Ile 40 Glu Gly Pro Ile Ala Leu Lys Phe Ser His Leu Cys Leu Glu Asp His 55 60 Asn Ser Tyr Cys Ile Asn Gly Ala Cys Ala Phe His His Glu Leu Glu 75 Lys Ala Ile Cys Arg Cys Phe Thr Gly Tyr Thr Gly Glu Arg Cys Leu 85 90 Lys Leu Lys Ser Pro Tyr Asn Val Cys Ser Gly Glu Arg Arg Pro Leu 105

<210> 1651 <211> 50 <212> PRT <213> Homo sapiens

<210> 1652 <211> 121 <212> PRT <213> Homo sapiens

 <400> 1652

 Met Ser Arg Ala Gly Met Leu Gly Val Val Cys Ala Leu Leu Val Trp 1

 1
 5
 10
 15

 Ala Tyr Leu Ala Val Gly Lys Leu Val Val Arg Met Thr Phe Thr Glu 20
 25
 30

 Leu Cys Thr His His Pro Trp Ser Leu Arg Cys Glu Ser Phe Cys Arg 35
 40
 45

 Ser Arg Val Thr Ala Cys Leu Pro Ala Pro Ala Pro Trp Leu Arg Pro 50
 55
 60

 Phe Leu Cys Pro Met Leu Phe Ser Asp Arg Asn Pro Val Glu Cys His 65
 70
 75
 80

 Leu Phe Gly Glu Ala Val Ser Asp Pro Val Cys Lys Gly Leu Pro 95
 95

 His Tyr Phe Trp His Pro Thr Phe Phe Pro Val Lys Ala Asn Cys Leu 100
 105
 110

 Val Ser Phe Cys Pro Thr Thr Val *
 110

<210> 1653 <211> 111 <212> PRT <213> Homo sapiens

100 105 110

<210> 1654 <211> 150 <212> PRT <213> Homo sapiens

<400> 1654 Met Trp Ile Cys Arg Val Lys Gln Ala Trp Leu Pro Pro Leu Leu Ser 10 Pro Leu Gly Pro Pro Thr Pro Trp Asp Pro Phe Tyr Ala Ala Pro Ser 25 Pro Pro Val Trp Val Gly Ser Gly Tyr Trp Tyr Arg Gly Leu Leu Ser 40 45 Pro Pro Asp Gly Gly Gln Gly Ser Phe Pro Pro His Leu Cys Pro Gln 55 Cys Pro Val Gln Ala Gln Ala Gln Ile Gly Pro Tyr Phe Arg Glu Leu 70 75 Gly Glu Pro Pro Ser Glu Thr Lys Trp Tyr Leu Asn Ser His Ser His 90 His Arg Ala Ala Gly Thr Gln Arg Arg Leu Arg Cys Leu Gln His Leu 105 Leu Gly Gly Gly Pro Gly Ile Gly Ser Glu Ser Pro Asn Glu Gly 120 Pro Gly Gln Val Thr His Ala Cys Asn Leu Ser Thr Leu Gly Gly Lys 135 Asp Val Arg Ile Thr * 145 149

<210> 1655 <211> 68 <212> PRT <213> Homo sapiens

<210> 1656 <211> 61 <212> PRT <213> Homo sapiens

<210> 1657 <211> 80 <212> PRT <213> Homo sapiens

<210> 1658 <211> 160 <212> PRT <213> Homo sapiens

<400> 1658

Met Ala Phe Leu Leu Tyr His Leu Val Tyr His Ile Pro Pro Met Ala 10 Pro Val Ser Phe Val Phe Glu Thr Lys Ser Arg Ser Ala Ala Gln Ala 25 Gly Val Gln Trp His Asp Pro Gly Ser Pro Gln Pro Leu Pro Pro Arg 40 Phe Lys Arg Phe Ser Cys His Gly Leu Asn Ile Lys Phe Ala Phe Phe 55 Ser His Leu Lys Glu Leu His Leu Asp Ser Gly His Cys Phe Ile Phe 75 70 Ile Arg Leu Val Lys Gly Ala Val Cys Leu Ile His Val Gln Ile Arg 90 85 Ile Pro Ser Ala Asp Glu Asp Ile Thr Ile Leu Phe Phe Ile Val Ser 105 Lys His Phe Leu Glu Ser Val Phe Lys Met Leu Gln Trp Ser Gln Met 125 120 Thr Leu Ala Thr Val Lys Thr Thr Phe Ile Gly Leu Asn Glu Phe Ile 135

Cys Ser Pro Ser Thr Leu Pro Ser Gly Lys Lys Asn Gly Leu Ile *

145 150 155 159

<210> 1659 <211> 90 <212> PRT <213> Homo sapiens

vars nome suprem

<400> 1659 Met Trp Arg Leu Pro His Ser Gln Phe Ile His Ile Val Ile Leu Pro 5 10 Leu Lys Val Phe Leu Phe Leu Phe Cys Phe Leu Arg Trp Ser Phe Ser 20 25 Leu Val Ala Gln Ala Gly Val Gln Trp Arg Asp Leu Gly Pro Leu Gln 40 45 Pro Pro Pro Pro Arg Leu Lys Arg Phe Phe Cys Leu Ser Leu Pro Ser 55 60 Ser Trp Asp Tyr Arg His Ser Pro Pro His Pro Ala Asn Phe Tyr Thr 70 Phe Gly Arg Asp Gly Val Ser Pro Cys * 85

<210> 1660 <211> 56 <212> PRT <213> Homo sapiens

<210> 1661 <211> 74 <212> PRT <213> Homo sapiens

Asp Gly Thr Glu Gly His Tyr Pro Lys * 65 70 73

<210> 1662 <211> 271 <212> PRT <213> Homo sapiens

<400> 1662 Met Gly Leu Gly Gln Pro Gln Ala Trp Leu Leu Gly Leu Pro Thr Ala 5 10 Val Val Tyr Gly Ser Leu Ala Leu Phe Thr Thr Ile Leu His Asn Val 25 Phe Leu Leu Tyr Tyr Val Asp Thr Phe Val Ser Val Tyr Lys Ile Asn 40 Lys Met Ala Phe Trp Val Gly Glu Thr Val Phe Leu Leu Trp Asn Ser 55 Leu Asn Asp Pro Leu Phe Gly Trp Leu Ser Asp Arg Gln Phe Leu Ser 70 Ser Gln Pro Arg Ser Gly Ala Gly Leu Ser Ser Arg Ala Val Leu 90 85 Ala Arg Val Gln Ala Leu Gly Trp His Gly Pro Leu Leu Ala Leu Ser 105 100 Phe Leu Ala Phe Trp Val Pro Trp Ala Pro Ala Gly Leu Gln Phe Leu 120 Leu Cys Leu Cys Leu Tyr Asp Gly Phe Leu Thr Leu Val Asp Leu His 135 140 His His Ala Leu Leu Ala Asp Leu Ala Leu Ser Ala His Asp Arg Thr 150 155 His Leu Asn Phe Tyr Cys Ser Leu Phe Ser Ala Ala Gly Ser Leu Ser 165 170 Val Phe Ala Ser Tyr Ala Phe Trp Asn Lys Glu Asp Phe Ser Ser Phe 180 185 Arg Ala Phe Cys Val Thr Leu Ala Val Ser Ser Gly Leu Gly Phe Leu 200 Gly Ala Thr Gln Leu Leu Arg Arg Arg Val Glu Ala Ala Arg Lys Asp 215 Pro Gly Cys Ser Gly Leu Val Val Asp Ser Gly Leu Cys Gly Glu Glu 235 Leu Leu Val Gly Ser Glu Glu Ala Asp Ser Ile Thr Leu Gly Arg Tyr 245 250 Leu Arg Gln Leu Ala Arg His Arg Asn Phe Leu Cys Phe Ser * 265

<210> 1663 <211> 53 <212> PRT <213> Homo sapiens

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20
                                25
Lys Tyr Asn Thr Ser Ser Glu Tyr Leu Ser Glu Leu Asp Thr Glu Ala
       35
                            40
Ser Arg Val Ser
     50
            52
     <210> 1664
     <211> 271
     <212> PRT
     <213> Homo sapiens
     <400> 1664
Met Gly Leu Gly Gln Pro Gln Ala Trp Leu Leu Gly Leu Pro Thr Ala
                                    10
Val Val Tyr Gly Ser Leu Ala Leu Phe Thr Thr Ile Leu His Asn Val
                                25
Phe Leu Leu Tyr Tyr Val Asp Thr Phe Val Ser Val Tyr Lys Ile Asn
                            40
Lys Met Ala Phe Trp Val Gly Glu Thr Val Phe Leu Leu Trp Asn Ser
                        55
Leu Asn Asp Pro Leu Phe Gly Trp Leu Ser Asp Arg Gln Phe Leu Ser
                                        75
Ser Gln Pro Arg Ser Gly Ala Gly Leu Ser Ser Arg Ala Val Leu
                85
                                    90
Ala Arg Val Gln Ala Leu Gly Trp His Gly Pro Leu Leu Ala Leu Ser
           100
                                105
Phe Leu Ala Phe Trp Val Pro Trp Ala Pro Ala Gly Leu Gln Phe Leu
                           120
Leu Cys Leu Cys Leu Tyr Asp Gly Phe Leu Thr Leu Val Asp Leu His
                       135
His His Ala Leu Leu Ala Asp Leu Ala Leu Ser Ala His Asp Arg Thr
                  150
His Leu Asn Phe Tyr Cys Ser Leu Phe Ser Ala Ala Gly Ser Leu Ser
               165
                                   170
Val Phe Ala Ser Tyr Ala Phe Trp Asn Lys Glu Asp Phe Ser Ser Phe
Arg Ala Phe Cys Val Thr Leu Ala Val Ser Ser Gly Leu Gly Phe Leu
Gly Ala Thr Gln Leu Leu Arg Arg Arg Val Glu Ala Ala Arg Lys Asp
                       215
Pro Gly Cys Ser Gly Leu Val Val Asp Ser Gly Leu Cys Gly Glu Glu
                   230
                                       235
Leu Leu Val Gly Ser Glu Glu Ala Asp Ser Ile Thr Leu Gly Arg Tyr
               245
                                   250
Leu Arg Gln Leu Ala Arg His Arg Asn Phe Leu Cys Phe Ser *
                               265
    <210> 1665
    <211> 284
    <212> PRT
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<400> 1665

<213> Homo sapiens

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Met Asp Glu Lys Ser Asn Lys Leu Leu Leu Ala Leu Val Met Leu Phe
                                    10
Leu Phe Ala Val Ile Val Leu Gln Tyr Val Cys Pro Gly Thr Glu Cys
            20
                                 25
Gln Leu Leu Arg Leu Gln Ala Phe Ser Ser Pro Val Pro Asp Pro Tyr
                            40
Arg Ser Glu Asp Glu Ser Ser Ala Arg Phe Val Pro Arg Tyr Asn Phe
                        55
                                             60
Thr Arg Gly Asp Leu Leu Arg Lys Val Asp Phe Asp Ile Lys Gly Asp
                     70
                                         75
Asp Leu Ile Val Phe Leu His Ile Gln Lys Thr Gly Gly Thr Thr Phe
                85
                                    90
Gly Arg His Leu Val Arg Asn Ile Gln Leu Glu Gln Pro Cys Glu Cys
                                105
           100
Arg Val Gly Gln Lys Lys Cys Thr Cys His Arg Pro Gly Lys Arg Glu
        115
                            120
Thr Trp Leu Phe Ser Arg Phe Ser Thr Gly Trp Ser Cys Gly Leu His
                        135
                                            140
Ala Asp Trp Thr Glu Leu Thr Ser Cys Val Pro Ser Val Gly Asp Gly
                    150
                                       155
Lys Arg Asp Ala Arg Leu Arg Pro Ser Arg Trp Arg Ile Phe His Ile
                165
                                    170
Leu Tyr Ala Ala Cys Thr Asp Ile Arg Gly Ser Pro Asn Thr Asn Ala
                                185
Gly Ala Asn Ser Pro Ser Phe Thr Lys Thr Arg Asn Thr Ser Lys Ser
                            200
Trp Lys Asn Phe His Tyr Ile Thr Ile Leu Gln Asp Pro Gly Ala Arg
                                    220
                        215
Ser Leu Ser Glu Trp Arg Pro Val Leu Lys Arg Gly Thr Leu Glu Gly
                                        235
                    230
Leu Leu Ala Cys Trp Pro Trp Lys Ala Pro Pro Pro Leu Lys Lys Leu
                245
                                    250
Ser Thr Trp Tyr Pro Gly Glu Glu Leu Val Trp Leu Ala Pro Leu Gln
                                265
Lys Ile Ile Gly Leu Ala Leu Leu Ile Tyr Pro
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<210> 1666 <211> 67 <212> PRT

<213> Homo sapiens

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<210> 1667 <211> 79 <212> PRT <213> Homo sapiens

<400> 1667

Met Asn Thr His Trp Asn Ile Leu Pro Val Glu Arg Ser Cys Pro Leu 10 Trp Ile Ser Ser Glu Leu Ser Tyr Cys Ser Ile Lys Leu Leu Phe Ile 20 25 Leu Leu Thr Leu His Leu Pro Ala Tyr Leu Ile Leu Pro Gly His Lys 35 40 45 Ile Arg Thr Gln Asp Leu Pro Asn Glu Ala Asn Arg Ala Val Thr Gln 55 60 Thr Gly Leu Arg His Ala Leu Tyr Gln Ser Ile Ser Cys Trp * 70

<210> 1668 <211> 54 <212> PRT <213> Homo sapiens

<400> 1668 Met Trp Gly Leu Leu Ile Pro Cys Ile Leu Gly Cys Met Lys Leu Pro 1 5 10 His Asn Leu Leu Met Leu Phe Ser Leu Glu Thr Phe Leu Thr Leu Arg 20 25 Phe Ile Leu Asp Ser Phe Tyr Ser Tyr Val Phe Lys Pro Thr Asn Lys 35 40 Arg Phe Cys Asn Ile * 53

<210> 1669 <211> 119

50

<212> PRT <213> Homo sapiens

<400> 1669 Met Met Ala Gly Ile Arg Ala Leu Phe Met Tyr Leu Trp Leu Gln Leu 10 Asp Trp Val Ser Arg Gly Glu Ser Val Gly Leu His Leu Pro Thr Leu 25 Ser Val Gln Glu Gly Asp Asn Ser Ile Ile Asn Cys Ala Tyr Ser Asn 40 Ser Ala Ser Asp Tyr Phe Ile Trp Tyr Lys Gln Glu Ser Gly Lys Gly . 60 55 Pro Gln Phe Ile Ile Asp Ile Arg Ser Asn Met Asp Lys Arg Gln Gly 70 75 Gln Arg Val Thr Val Leu Leu Asn Lys Thr Val Lys His Leu Ser Leu 85 90 Gln Ile Ala Ala Thr Gln Pro Gly Asp Ser Ala Val Tyr Phe Cys Ala 105

Glu Ile Pro Glu Gln Arg * 115 118

<210> 1670

<211> 116

<212> PRT

<213> Homo sapiens

<400> 1670

Met Cys Leu Cys Cys Glu Cys Leu Phe His Leu Trp Lys Arg Ile 10 Asn Trp Trp Gln Gly Phe Cys Ser Phe Tyr Leu Leu Trp Val Gly 25 Leu Leu Ser Phe Pro Pro Asp Pro Pro Trp Lys Ser Phe Thr Pro Ala 40 Ile Leu Phe Leu Ala Trp Gly Thr Gly Ser Ser Pro Gly Arg His Arg 55 60 Phe Ser Leu Pro Thr Asp Arg Pro Ser Ala His Ser Pro Phe Leu 70 75 Ser Thr Leu Gln His Ser Ile Arg Thr Leu Phe His Ser Pro Ile Arg 85 90 Ser Ser Arg Phe Ala Phe Val Ser Ser Leu His Ser Tyr Thr Ser Ile 105 Pro Ser Leu Pro

<210> 1671

115 116

<211> 70

<212> PRT

<213> Homo sapiens

<400> 1671

 Met
 Ser
 His
 Cys
 Gly
 Leu
 Leu
 Phe
 Leu
 Val
 Thr
 Trp
 Leu
 Leu
 Ser

 Phe
 Ile
 Phe
 Leu
 Val
 Cys
 Lys
 Met
 Arg
 Ile
 Thr
 Phe
 Leu
 Phe
 Cys
 Leu

 Leu
 Thr
 Val
 Asp
 Met
 Lys
 Pro
 Asp
 Lys
 Val
 Leu
 Tyr
 Met
 Lys
 Cys
 Phe

 Lys
 Cys
 Ile
 Leu
 Leu
 Ser
 Cys
 Tyr
 Pro
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 Val
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 Lys
 Asp
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 Glu
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<210> 1672

<211> 263

<212> PRT

<213> Homo sapiens

<400> 1672

Met Arg Val Leu Cys Ala Phe Pro Glu Ala Met Pro Ser Ser Asn Ser

10 Arg Pro Pro Ala Cys Leu Ala Pro Gly Ala Leu Tyr Leu Ala Leu Leu 25 Leu His Leu Ser Leu Ser Ser Gln Ala Gly Asp Arg Pro Leu Pro 40 Val Asp Arg Ala Ala Gly Leu Lys Glu Lys Thr Leu Ile Leu Leu Asp 55 Val Ser Thr Lys Asn Pro Val Arg Thr Val Asn Glu Asn Phe Leu Ser 70 75 Leu Gln Leu Asp Pro Ser Ile Ile His Asp Gly Trp Leu Asp Phe Leu 85 90 Ser Ser Lys Arg Leu Val Thr Leu Ala Arg Gly Leu Ser Pro Ala Phe 1.00 105 Leu Arg Phe Gly Gly Lys Arg Thr Asp Phe Leu Gln Phe Gln Asn Leu 120 125 Arg Asn Pro Ala Lys Ser Arg Gly Gly Pro Gly Pro Asp Tyr Tyr Leu 135 140 Lys Asn Tyr Glu Asp Asp Ile Val Arg Ser Asp Val Ala Leu Asp Lys 150 155 Gln Lys Gly Cys Lys Ile Ala Gln His Pro Asp Gly Met Leu Glu Pro 170 Pro Arg Glu Lys Ala Ala Gln Met His Leu Val Leu Lys Glu Gln 180 185 Phe Ser Asn Thr Tyr Ser Asn Leu Ile Leu Thr Glu Pro Asn Asn Tyr 200 Arg Thr Met His Gly Arg Ala Val Asn Gly Ser Gln Leu Gly Lys Asp 215 Tyr Ile Gln Leu Lys Ser Leu Leu Gln Pro Ile Arg Ile Tyr Ser Arg 230 Ala Ser Leu Tyr Gly Pro Asn Ile Val Arg Pro Arg Lys Asn Val Ile 245 250 Ala Leu Leu Asp Gly Leu * 260 262

<210> 1673 <211> 156 <212> PRT

<213> Homo sapiens

<400> 1673

 Met
 Lys
 Trp
 Lys
 Thr
 Gly
 Val
 Ala
 Ile
 Phe
 Val
 Val
 Val
 Val
 Tyr

 Leu
 Val
 Thr
 Gly
 Leu
 Val
 Phe
 Arg
 Ala
 Leu
 Glu
 Glu
 Pro
 Phe
 Glu

 Ser
 Ser
 Gln
 Lys
 Asn
 Thr
 Ile
 Ala
 Leu
 Glu
 Lys
 Ala
 Glu
 Phe
 Leu
 Arg

 Asp
 His
 Val
 Ser
 Pro
 Glu
 Leu
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 Thr
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 Ile
 Glu
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Phe Gly Phe Leu Leu Ala Gly Ile Glu Asp Gln Leu Gly Thr Ile Phe 130 135 140 Gly Lys Ser Ile Ala Arg Val Glu Lys Val Phe * 145 150 155

<210> 1674 <211> 83 <212> PRT <213> Homo sapiens

<210> 1675 <211> 54 <212> PRT <213> Homo sapiens

<210> 1676 <211> 119 <212> PRT <213> Homo sapiens

<210> 1677 <211> 49 <212> PRT <213> Homo sapiens

<210> 1678 <211> 127 <212> PRT <213> Homo sapiens

<210> 1679

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<211> 49
     <212> PRT
     <213> Homo sapiens
     <400> 1679
Met Ile Phe Phe Ile Lys Ala Pro Leu Tyr Leu Leu Gln Ser Met Met
Asp Cys Leu Tyr Ala Arg Ile Pro Cys Ile Thr Asp Cys Ala Met
            20
                                25
Ala Glu Ile Glu Lys Leu Gly Gln Lys Tyr Pro Val Ala Leu Arg Ile
                            40
Ala
49
     <210> 1680
     <211> 58
     <212> PRT
     <213> Homo sapiens
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<400> 1680 Met Val Tyr Glu Val Phe Ile Asn Lys Ala Asn Ile Leu Leu Leu 10 Phe Leu Arg Gln Ser Leu Ala Val Leu Pro Arg Leu Glu Cys Ser Gly 25 Ala Ile Ser Ala Arg Cys Asn Leu His Leu Arg Ile Pro Pro Asp Phe 40

His Arg Ser Thr Met Gly Gly Gly Gly 55

> <210> 1681 <211> 49 <212> PRT <213> Homo sapiens

<400> 1681 Met Leu Ser Gly Trp Val Gln Cys Pro Leu Leu Gln Arg Val His Phe 10 Tyr Ala Phe Ser Val Gly Pro Phe His Arg Lys Ile Trp Gly Asp Val 25 Ser Phe Pro Leu Thr Phe Tyr Phe Lys Asn Leu Gln Thr Gln Lys Ser 35 40 45

<210> 1682 <211> 78 <212> PRT <213> Homo sapiens . .

| Met | Thr | Gly | Leu | Phe | Leu | His | His | Asn | Pro | Gly | Ile | Leu | Leu | Ala | Pro | Ile | Val | Leu | Leu | Ala | Pro | Ile | Val | Leu | Leu | Ala | Pro | Ile | Val | Leu | Ala | Pro | Ile | Val | Leu | Ala | Pro | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile | Ile

<210> 1683 <211> 52 <212> PRT <213> Homo sapiens

<210> 1684 <211> 165 <212> PRT <213> Homo sapiens

<400> 1684 Met Pro Ala Pro Pro Leu Pro Gly Gly Trp Asn Thr Trp Gly Pro Ser 1 5 10 Leu Ser Leu Pro Leu Leu Leu Gly Ala Val Ala Met Ala Leu Gly 25 Val Arg Pro Pro Gly Gln Val Gly Leu Ser Pro Ile Ala Thr Ala Ser 40 Thr Val Gly Val Pro Arg Cys Leu Gln Thr Ala Phe Arg Gly Asp Ala 55 Gly Trp His Ser Cys Ala Gln Gln Gly Ala Cys Val Ala Leu His Pro 75 Ser Glu Arg Arg Leu Gly Ile Ser Asp Glu Ala His Ser Arg Ser Arg . 90 Trp Gly Gly Glu Asp Ser Pro Ser Pro Leu Thr Gly Pro Pro Leu Ser 105 110 Pro Ser Pro Pro Asp Cys Leu Ser Leu Pro Arg Leu Thr Pro Leu Arg 120 Leu Pro Pro Pro Phe Pro Phe Leu Gly Pro Ile Pro Ser Leu Pro 135 140 Pro Pro Pro Ser Pro Pro Pro Gln Pro Pro Ala Thr Ala Pro Pro 150 155

Ser Leu Arg Phe * 164

<210> 1685 <211> 153 <212> PRT <213> Homo sapiens

<400> 1685 Met Gly Thr Ala Ala Leu Gly Pro Val Trp Ala Ala Leu Leu Phe 10 Leu Leu Met Cys Glu Ile Pro Met Val Glu Leu Thr Phe Asp Arg Ala 25 Val Ala Ser Gly Cys Gln Arg Cys Cys Asp Ser Glu Asp Pro Leu Asp 40 Pro Ala His Val Ser Ser Ala Ser Ser Ser Gly Arg Pro His Ala Leu 55 Pro Glu Ile Arg Pro Tyr Ile Asn Ile Thr Ile Leu Lys Ala Gln Arg 70 75 Ala Gln His His Ala Glu Pro Glu Cys Asp Ala Gly Pro Gly Leu Arg 85 90 Gly Pro Arg Leu Gly Ala Ala Leu Gln Ala Pro Ala Arg Glu Arg His 100 105 Leu Gln Gln Arg Leu Arg His Leu His His Leu Gln Arg Pro Pro His 120 125 Gln Gly Arg Gly Arg Leu Arg Ala Ser Gly Pro Pro Ser Arg Leu Glu 135 Ser Ser Ala Asp Pro Ala Pro Ala * 145 150

<210> 1686 <211> 141 <212> PRT <213> Homo sapiens

<400> 1686 Met Arg Arg Thr Ala Phe Ile Leu Gly Ser Gly Leu Leu Ser Phe Val 10 Ala Phe Trp Asn Ser Val Thr Trp His Leu Gln Arg Phe Trp Gly Ala 25 Ser Gly Tyr Phe Trp Gln Ala Gln Trp Glu Arg Leu Leu Thr Thr Phe 40 Glu Gly Lys Glu Trp Ile Leu Phe Phe Ile Gly Ala Ile Gln Val Pro 55 Cys Leu Phe Phe Trp Ser Phe Asn Gly Leu Leu Leu Val Val Asp Thr 70 Thr Gly Lys Pro Asn Phe Ile Ser Arg Tyr Arg Ile Gln Val Gly Lys 90 Asn Glu Pro Val Asp Pro Val Lys Leu Arg Gln Ser Ile Arg Thr Val 105 Leu Phe Asn Gln Cys Met Ile Ser Phe Pro Met Gly Gly Leu Pro Leu 120 Ser Leu Pro Gln Met Val Glu Arg Pro Leu Thr Pro *

130 135 140

<210> 1687 <211> 61 <212> PRT

<213> Homo sapiens

Pro Trp Val Gly Cys Gly Arg Gly Glu Asn Pro Ser Val Gly Thr Val

Asp Pro Ser Cys Arg Leu Cys Ala Pro Gly His Val * 50 55 60

<210> 1688

<211> 68

<212> PRT

<213> Homo sapiens

<400> 1688

 Met
 Val
 Ala
 Ala
 Thr
 Pro
 Pro
 Gly
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 Trp
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 Leu
 Val
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<210> 1689

67

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<211> 74

<212> PRT

<213> Homo sapiens

<400> 1689

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<210> 1690
     <211> 114
     <212> PRT
     <213> Homo sapiens
    <400> 1690
Met His Met Cys Ala Phe Leu His Val Trp Thr Cys Ala Cys Met His
                 5
                                     10
Leu Cys Val Cys Val Cys Ala Glu Thr Gly Lys Gly Val Lys Val Leu
            20
                                 25
Val Arg Glu Pro Gly Ser Phe Leu Phe Pro Asn Leu Ser Cys Ser Lys
                             40
Glu Gly Trp Gly Trp Gly Gln Pro Leu Leu Lys Val Ile Gly Glu Glu
                         55
Arg Phe Ser Asp Ser Glu Val Thr Ala Ser Val Ala Gln Ala Val Ser
                     70
                                         75
Leu Val Thr Val Ile Leu Gln Phe Ser Asp Pro His Val Ser Phe Arg
                                     90
                 85
Gly Lys Arg Lys Lys Gly Thr Leu Trp Trp Val Leu Gly Gly Lys Arg
            100
                                105
Lys *
113
     <210> 1691
     <211> 69
     <212> PRT
     <213> Homo sapiens
     <400> 1691
Met Ala Phe Leu Leu Ser Thr Leu Leu Asn His Tyr Leu Ala Cys Lys
His Ser Ser Glu Leu Trp Leu Gln Ser Ser Leu Asn Asn Leu Gly Lys
                                 25
Lys Lys Asp Lys Ala Tyr Ile Phe Thr Val Leu Ala Leu Lys His Ile
                            40
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<210> 1692 <211> 103 <212> PRT <213> Homo sapiens

69

<400> 1692

Leu Met Pro Val Ser

Met Leu Gly Pro Thr Val Phe Asn Ile Lys Phe Val Phe Leu Ile Thr 1 5 10 15 Ala Leu Gly Ala Leu Pro Ser Ser Leu Pro His Ala His Ser Ala Ala

Pro Gln Met Pro Leu Arg Ile Tyr Phe Val Leu Gly Gln Ser Trp Trp

<210> 1693 <211> 48 <212> PRT <213> Homo sapiens

<210> 1694 <211> 92 <212> PRT <213> Homo sapiens

<210> 1695 <211> 83 <212> PRT <213> Homo sapiens

<210> 1696 <211> 159 <212> PRT <213> Homo sapiens

<400> 1696 Met Leu Trp Leu Phe Gln Ser Leu Leu Phe Val Phe Cys Phe Gly Pro 5 10 Gly Asn Val Val Ser Gln Ser Ser Leu Thr Pro Leu Met Val Asn Gly 2.0 25 Ile Leu Gly Glu Ser Val Thr Leu Pro Leu Glu Phe Pro Ala Gly Glu 40 Lys Val Asn Phe Ile Thr Trp Leu Phe Asn Glu Thr Ser Leu Ala Phe 55 Ile Val Pro His Glu Thr Lys Ser Pro Glu Ile His Val Thr Asn Pro 70 Lys Gln Gly Lys Arg Leu Asn Phe Thr Gln Ser Tyr Ser Leu Gln Leu 90 Ser Asn Leu Lys Met Glu Asp Thr Gly Ser Tyr Arg Ala Gln Ile Ser 105 Thr Lys Thr Ser Ala Lys Leu Ser Ser Tyr Thr Leu Arg Ile Leu Thr 120 125 Leu Tyr Pro Ile Val Gly Asn Gly Ile Trp Gly Asn Lys Asn Phe Leu 135 140 Thr Thr Leu Ala Arg Gly Asn Val Lys Leu Asp Gly Leu His Glu 150 155

<210> 1697 <211> 105 <212> PRT <213> Homo sapiens

 Pro
 Gln
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 Thr
 Thr
 Val
 Leu
 Asp
 Leu
 Arg
 Phe
 Asn
 Arg
 Ile
 Arg
 Glu

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 Gly
 Ser
 Ala
 Phe
 Lys
 Lys
 Asn
 Leu
 Asn
 Thr
 Leu
 Tyr

 65
 Tyr
 Lys
 Asn
 Glu
 His
 Ala
 Leu
 Asp
 Lys
 Gln
 Thr
 Phe
 Lys
 Gly

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 Lys
 Asn
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<210> 1698 <211> 195 <212> PRT <213> Homo sapiens

<400> 1698 Met Pro Ser Trp Ile Gly Ala Val Ile Leu Pro Leu Leu Gly Leu Leu 5 10 Leu Ser Leu Pro Ala Gly Ala Asp Val Lys Ala Arg Ser Cys Gly Glu 25 Val Arg Gln Ala Tyr Gly Ala Lys Gly Phe Ser Leu Ala Asp Ile Pro 40 Tyr Gln Glu Ile Ala Gly Glu His Leu Arg Ile Cys Pro Gln Glu Tyr 55 60 Thr Cys Cys Thr Thr Glu Met Glu Asp Lys Leu Ser Gln Gln Ser Lys 70 75 Leu Glu Phe Glu Asn Leu Val Glu Glu Thr Ser His Phe Val Arg Thr 90 Thr Phe Val Ser Arg His Lys Lys Phe Asp Glu Phe Phe Arg Glu Leu 105 Leu Glu Asn Ala Glu Lys Ser Leu Asn Asp Met Phe Val Arg Thr Tyr 120 Gly Met Leu Tyr Met Gln Asn Ser Glu Val Phe Gln Asp Leu Phe Thr 135 Glu Leu Lys Arg Tyr Tyr Thr Gly Gly Asn Val Asn Leu Glu Glu Met 150 155 Leu Asn Asp Phe Trp Ala Arg Leu Leu Glu Arg Met Phe Gln Leu Ile 165 770 Asn Pro Gln Tyr Pro Phe Ser Glu Gly Phe Leu Gly Met Cys Glu Gln 185 Ile Pro * 194

<210> 1699 <211> 97 <212> PRT <213> Homo sapiens

<400> 1699
Met Asp Ser Pro Trp Ala Gly Leu Leu Trp Leu Leu Pro Thr Leu Trp
1 5 10 15
Ser Ser Phe Pro Ala Pro Ala Cys Trp Pro Ser Ser Ser Ser Ser Ser
20 25 30

 Pro
 Val
 Cys
 Ala
 Ala
 Asn
 Gly
 Ala
 Met
 Ser
 Ala
 Ser
 Arg
 Asn
 Leu
 Arg

 Thr
 Leu
 Lys
 Gly
 Arg
 Thr
 Ala
 Pro
 Gly
 Ser
 Thr
 Leu
 Pro
 Leu
 Arg
 Arg
 Arg
 Arg
 Arg
 Arg
 Cys
 Leu
 Met
 Ser
 Thr
 Phe
 Ser
 Arg
 Trp

 65
 70
 75
 75
 80

 Leu
 Arg
 Ser
 Leu
 Pro
 Arg
 Ser
 Leu
 His
 Thr
 Gln
 Thr

 85
 90
 95
 96

<210> 1700 <211> 129 <212> PRT <213> Homo sapiens

<400> 1700 Met Gly Trp Ala Pro Leu Leu Thr Leu Leu Ala His Cys Thr Gly 10 Ser Trp Ala Gln Ser Val Leu Thr Gln Pro Pro Ser Glu Ser Glu Ala 20 25 Pro Gly Gln Trp Val Asn Ile Ser Cys Thr Gly Ser Gly Ser Asn Leu 40 Gly Ala Gly Phe Asp Val Gln Trp Tyr Gln Leu Ile Pro Gly Thr Ala 55 Pro Lys Leu Leu Ile Phe Asn Asn Asn Arg Gln Pro Ser Gly Val Pro 70 75 Asp Arg Phe Ser Ala Ser Lys Ser Gly Thr Ser Ala Ser Leu Thr Ile 8.5 90 Asn Asp Leu Gln Pro Glu Asp Glu Ser Glu Tyr Tyr Cys Leu Ala Met 105 Thr Ala Ala Ser Leu Val Ser Ser Glu Leu Gly Pro Lys Ser Pro Ala 115 120 125

<210> 1701 <211> 219 <212> PRT <213> Homo sapiens

85 90 Arg Trp Asn Glu Ile Phe Gly Asn Asn Leu Gly Ala Leu Ala Met Phe 105 100 Cys Val Leu Tyr Pro Glu Asn Ile Glu Ala Arg Asp Met Ala Lys Asp 120 125 Tyr Met Glu Arg Met Ala Ala Gln Pro Ser Trp Leu Val Lys Asp Ala 135 Pro Trp Asp Glu Val Pro Leu Ala His Ser Leu Val Gly Phe Ala Thr 150 155 Ala Tyr Asp Phe Leu Tyr Asn His Leu Ser Lys Thr Gln Gln Glu Lys 165 170 Phe Leu Glu Val Ile Ala Asn Ala Ser Gly Tyr Met Phe Val Thr Leu 190 180 185 Ile Leu Gly Ala Asp Gly Asp Ser Asn Thr Cys Thr Ile Ile Ser Pro 200 205 Pro Thr Val Trp Leu Cys Ser Arg Glu Ala * 215

<210> 1702 <211> 86 <212> PRT <213> Homo sapiens

<210> 1703 <211> 229 <212> PRT <213> Homo sapiens

Phe Cys Asp Met Thr Ser Gly Gly Gly Gly Trp Thr Leu Val Ala Ser 85 Val His Glu Asn Asp Met His Gly Lys Cys Thr Val Gly Asp Arg Trp 100 105 Ser Ser Gln Gln Gly Asn Lys Ala Asp Tyr Pro Glu Gly Asp Gly Asn 120 125 Trp Ala Asn Tyr Asn Thr Phe Gly Ser Ala Glu Ala Ala Thr Ser Asp 135 140 Asp Tyr Lys Asn Pro Gly Tyr Tyr Asp Ile Gln Ala Lys Asp Leu Gly 150 155 Ile Trp His Val Pro Asn Lys Ser Pro Met Gln His Trp Arg Asn Ser 165 170 Ala Leu Leu Arg Tyr Arg Thr Asn Thr Gly Phe Leu Gln Arg Leu Gly 180 185 His Asn Leu Phe Gly Ile Tyr Gln Lys Tyr Pro Val Lys Tyr Arg Ser 200 205 Gly Lys Cys Trp Asn Asp Asn Gly Pro Ala Ile Pro Trp Val Tyr Asp 215 Phe Gly Glu Ala * 228

<210> 1704

<211> 202

<212> PRT

<213> Homo sapiens

<400> 1704 Met Val Phe Pro Val Met Tyr Asn Leu Ile Ile Leu Val Cys Arg Ala 10 Cys Phe Pro Asp Leu Gln His Gly Tyr Leu Val Ala Trp Leu Val Leu 20 25 Asp Tyr Thr Ser Asp Leu Leu Tyr Leu Leu Asp Met Val Val Arg Phe 40 His Thr Gly Phe Leu Glu Gln Gly Ile Leu Val Val Asp Lys Gly Arg Ile Ser Ser Arg Tyr Val Arg Thr Trp Ser Phe Phe Leu Asp Leu Ala Ser Leu Met Pro Thr Asp Val Val Tyr Val Arg Leu Gly Pro His Thr 90 Pro Thr Leu Arg Leu Asn Arg Phe Leu Arg Ala Pro Arg Leu Phe Glu 105 Ala Phe Asp Arg Thr Glu Thr Arg Thr Ala Tyr Pro Asn Ala Phe Cys 120 Ile Gly Lys Leu Met Leu Tyr Ile Phe Gly Arg Ile His Trp Asn Asn 135 140 Cys Leu Tyr Phe Ser Leu Ser Arg Tyr Leu Gly Phe Gly Arg Glu Pro 150 155 Met Gly Val Pro Arg Thr Pro Ala Pro Thr Trp Val Leu Thr Ala Arg 170 165 Gly Gly Pro Val Thr Ser Tyr Lys Leu Phe Asn Phe Phe His Pro Leu 180 185 Asp Thr Trp Ile Ile Gln Gly Gly Glu *

<210> 1705 <211> 58 <212> PRT <213> Homo sapiens

<400> 1705

 Met Gly Leu Leu Gly Val Leu Trp Asn Thr Thr Leu His Met Cys Arg

 1
 5
 10
 15

 Met Arg Leu Gln Asp Thr Gly Gln Lys Ile Arg Thr Gly Ser Cys Glu
 20
 25
 30

 Leu His Gly Ser Gln Ser Ser His Ser Thr Gly Asn Leu Arg Val Leu
 35
 40
 45

 Pro Ser His Asn Gly Glu Thr Leu His
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 50
 55
 57

<210> 1706 <211> 55 <212> PRT <213> Homo sapiens

<210> 1707 <211> 139 <212> PRT <213> Homo sapiens

<400> 1707 Met Leu Glu Cys Ala Phe Ile Val Leu Trp Leu Gln Leu Gly Trp Leu 10 Ser Gly Glu Asp Gln Val Thr Gln Ser Pro Glu Ala Leu Arg Leu Gln 25 Glu Gly Glu Ser Ser Ser Leu Asn Cys Ser Tyr Thr Val Ser Gly Leu 40 Arg Gly Leu Phe Trp Tyr Arg Gln Asp Pro Gly Lys Gly Pro Glu Phe 55 Leu Phe Thr Leu Tyr Ser Ala Gly Glu Glu Lys Glu Lys Glu Arg Leu 7.5 Lys Ala Thr Leu Thr Lys Lys Glu Ser Phe Leu His Ile Thr Ala Pro 90 Lys Pro Glu Asp Ser Ala Thr Tyr Leu Cys Ala Val Gln Ala Gln Phe 105 His Ser Gly Gly Gly Ala Asp Gly Leu Thr Phe Gly Lys Gly Thr Arg 120

Leu Lys Val Leu Ala Leu Tyr Pro Glu Pro * 130 135 138

<210> 1708

<211> 59

<212> PRT

<213> Homo sapiens

<400> 1708

 Met Gly Pro Arg
 Phe Val Ser Thr Leu Pro Phe Ser Pro Ser Ala Ala

 1
 5
 10
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 Trp Cys Ala Cys Glu Ala Gly Gly Gly Leu Arg Arg Glu Val Ala His
 20
 25
 30

 Ala Gln Arg Ala Ala Ser Thr Ala Pro Thr Ala His Met Gln Asn Ser
 45

 Thr Leu Ile Gly Leu Asn Leu Ser Arg Gly *
 55
 58

<210> 1709

<211> 81

<212> PRT

<213> Homo sapiens

<400> 1709

 Met
 Arg
 Leu
 Pro
 Trp
 Glu
 Leu
 Leu
 Val
 Leu
 Gln
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 Phe
 Ile
 Leu
 Cys

 Leu
 Ala
 Asp
 Asp
 Ser
 Thr
 Leu
 His
 Gly
 Pro
 Ile
 Phe
 Ile
 Glu
 Pro

 Ser
 Pro
 Val
 Met
 Phe
 Pro
 Leu
 Asp
 Ser
 Glu
 Glu
 Lys
 Ala
 Lys
 Leu

 Asn
 Cys
 Glu
 Asp
 Ala
 Asp
 Phr
 Glu
 Phe
 Leu
 Glu
 Arg
 Cys

 Asn
 Gly
 Ala
 Asp
 Thr
 Gly
 Met
 Glu
 Phe
 Leu
 Gln
 Arg
 Cys

 65
 70
 75
 80

<210> 1710

<211> 399

<212> PRT

<213> Homo sapiens

<400> 1710

 Met
 Leu
 Arg
 Leu
 Tyr
 Val
 Met
 Gly
 Val
 Ser
 Ala
 Phe
 Thr
 Leu

 1
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 Gln
 Pro
 Ala
 Ala
 His
 Thr
 Gly
 Ala
 Ala
 Arg
 Ser
 Cys
 Arg
 Phe
 Arg
 Gly
 Arg
 Gly
 Fro
 Val
 Ala
 Leu
 His
 Fro
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 Ala
 Leu
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 Gly
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55
Ile Asn Leu Thr Trp His Lys Asn Asp Ser Ala Arg Thr Val Pro Gly
                   70
Glu Glu Glu Thr Arg Met Trp Ala Gln Asp Gly Ala Leu Trp Leu Leu
                                   90
               8.5
Pro Ala Leu Gln Glu Asp Ser Gly Thr Tyr Val Cys Thr Thr Arg Asn
           100
                              105
Ala Ser Tyr Cys Asp Lys Met Ser Ile Glu Leu Arg Val Phe Glu Asn
                          120
Thr Asp Ala Phe Leu Pro Phe Ile Ser Tyr Pro Gln Ile Leu Thr Leu
                      135
                                         140
Ser Thr Ser Gly Val Leu Val Cys Pro Asp Leu Ser Glu Phe Thr Arg
                                     155
                  150
Asp Lys Thr Asp Val Lys Ile Gln Trp Tyr Lys Asp Ser Leu Leu Leu
             165
                                 170
Asp Lys Asp Asn Glu Lys Phe Leu Ser Val Arg Gly Thr Thr His Leu
          180
                              185
Leu Val His Asp Val Ala Leu Glu Asp Ala Gly Tyr Tyr Arg Cys Val
                         200
                                             205
Leu Thr Phe Ala His Glu Gly Gln Gln Tyr Asn Ile Thr Arg Ser Ile
210 215
                               220
Glu Leu Arg Ile Lys Lys Lys Glu Glu Thr Ile Pro Val Ile Ile
                 230
                                     235
Ser Pro Leu Lys Thr Ile Ser Ala Ser Leu Gly Ser Arg Leu Thr Ile
                                  250
Pro Cys Lys Val Phe Leu Gly Thr Gly Thr Pro Leu Thr Thr Met Leu
                              265
Trp Trp Thr Ala Asn Asp Thr His Ile Glu Ser Ala Tyr Pro Gly Gly
                         280
Arg Val Thr Glu Gly Pro Arg Gln Glu Tyr Ser Glu Asn Asn Glu Asn
                     295
Tyr Ile Glu Val Pro Leu Ile Phe Asp Pro Val Thr Arg Glu Asp Leu
                  310
                                      315
His Met Asp Phe Lys Cys Val Val His Asn Thr Leu Ser Phe Gln Thr
               325
                                  330
Leu Arg Thr Thr Val Lys Glu Ala Ser Ser Thr Phe Ser Trp Gly Ile
                              345
Val Leu Ala Pro Leu Ser Leu Ala Phe Leu Val Leu Gly Gly Ile Trp
       355 . 360
Met His Arg Arg Cys Lys His Arg Thr Gly Lys Ala Asp Gly Leu Thr
                      375
Val Leu Trp Pro His His Gln Asp Phe Gln Ser Tyr Pro Lys *
                  390
                                      395
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<210> 1711

<211> 254

<212> PRT

<213> Homo sapiens

<400> 1711

Ile Ser Cys Pro His Glu Cys Phe Glu Ala Ile Leu Ser Leu Asp Thr 55 Gly Tyr Arg Ala Pro Val Thr Leu Val Arg Lys Gly Cys Trp Thr Gly Pro Pro Ala Gly Gln Thr Gln Ser Asn Ala Asp Ala Leu Pro Pro Asp 85 90 Tyr Ser Val Val Arg Gly Cys Thr Thr Asp Lys Cys Asn Ala His Leu 100 105 Met Thr His Asp Ala Leu Pro Asn Leu Ser Gln Ala Pro Asp Pro Pro 120 125 Thr Leu Ser Gly Leu Glu Cys Tyr Ala Cys Ile Gly Val His Gln Asp 135 Asp Cys Ala Ile Gly Arg Ser Arg Arg Val Gln Cys His Gln Asp Gln 150 155 Thr Ala Cys Phe Gln Gly Asn Gly Arg Met Thr Val Gly Asn Phe Ser 165 170 Val Pro Val Tyr Ile Arg Thr Cys His Arg Ala Leu Leu His His Leu 185 180 Met Gly Thr Thr Ser Pro Trp Thr Ala Ile Gly Pro Pro Arg Gly Ser . 205 1.95 200 Cys Cys Glu Gly Tyr Leu Cys Asn Arg Lys Ser Met Thr Gln Pro Phe 215 220 Thr Ser Ala Ser Ala Thr Thr Pro Pro Arg Ala Leu Gln Val Leu Ala 230 235 Leu Leu Pro Val Leu Leu Val Gly Leu Ser Ala * 245 250

<210> 1712 <211> 124 <212> PRT

<213> Homo sapiens

<400> 1712 Met Thr Trp Leu Leu Val Ala Tyr Ala Asp Phe Val Val Thr Phe Val 10 Met Leu Leu Pro Ser Lys Asp Phe Trp Tyr Ser Val Val Asn Gly Val 25 Ile Phe Asn Cys Leu Ala Val Leu Ala Leu Ser Ser His Leu Arg Thr 40 Met Leu Thr Asp Pro Glu Lys Ser Ser Asp Cys Arg Pro Ser Ala Cys 55 Thr Val Lys Thr Gly Leu Asp Pro Thr Leu Val Gly Ile Cys Gly Glu 75 70 Gly Thr Glu Ser Val Gln Ser Leu Leu Gly Ala Val Pro Lys Gly 85 90 Asn Ala Thr Lys Glu Tyr Met Asp Glu Leu Ala Ala Glu Ala Arg Gly 105 Ser His Leu Gln Val Pro Gln Val Leu Leu Tyr * 115 120 123

<210> 1713

<211> 214

<212> PRT

<213> Homo sapiens

<400> 1713 Met Leu His Leu Val Phe Ile Leu Pro Ser Leu Met Leu Leu Ile Pro 10 His Ile Leu Leu Glu Asn Phe Ala Ala Ile Pro Gly His Arg Cys 20 25 Trp Val His Met Leu Asp Asn Asn Thr Gly Ser Gly Asn Glu Thr Gly 40 Ile Leu Ser Glu Asp Ala Leu Leu Arg Ile Ser Ile Pro Leu Asp Ser 55 Asn Leu Arg Pro Glu Lys Cys Arg Arg Phe Val His Pro Gln Trp Gln 70 75 Leu Leu His Leu Asn Gly Thr Ile His Ser Thr Ser Glu Ala Asp Thr 90 Glu Pro Cys Val Asp Gly Trp Val Tyr Asp Gln Ser Tyr Phe Pro Ser 100 105 Thr Ile Val Thr Lys Trp Asp Leu Val Cys Asp Tyr Gln Ser Leu Lys 120 125 Ser Val Val Gln Phe Leu Leu Leu Thr Gly Met Leu Val Gly Gly Ile 135 140 Ile Gly Gly His Val Ser Asp Arg Trp Leu Val Glu Ser Ala Arg Trp 150 155 Leu Ile Ile Thr Asn Lys Leu Asp Glu Gly Leu Lys Ala Leu Arg Lys 165 170 Val Ala Arg Thr Asn Gly Ile Lys Asn Ala Glu Arg Asn Pro Glu His 180 185 Arg Gly Cys Lys Ile His His Ala Gly Gly Ala Gly Cys Ser Thr Asp 195 200 Gln Asn Tyr Cys Val * 210 213

<210> 1714 <211> 178 <212> PRT

<213> Homo sapiens

<400> 1714 Met Ala Ala Ser Trp Ser Leu Leu Val Thr Leu Arg Pro Leu Ala Gln 5 Ser Pro Leu Arg Gly Arg Cys Val Gly Cys Gly Ala Trp Ala Ala Ala 20 25 Leu Ala Pro Leu Ala Thr Ala Pro Gly Lys Pro Phe Trp Lys Ala Tyr 40 Thr Val Gln Thr Ser Glu Ser Met Thr Pro Thr Ala Thr Ser Glu Thr 55 Tyr Leu Lys Ala Leu Ala Val Cys His Gly Pro Leu Asp His Tyr Asp 70 Phe Leu Ile Lys Ala His Glu Leu Lys Asp Asp Glu His Gln Arg Arg Val Ile Gln Cys Leu Gln Lys Leu His Glu Asp Leu Lys Gly Tyr Asn Ile Glu Ala Glu Gly Leu Phe Phe Lys Ala Phe Phe Lys Glu Gln Thr 120 125 Ser Lys Gly Pro Val Cys Leu Trp Arg Cys Trp Tyr Arg Lys Asn Asn

<210> 1715 <211> 76 <212> PRT <213> Homo sapiens

<210> 1716 <211> 83 <212> PRT <213> Homo sapiens

<400> 1716 Met Arg Phe Thr Phe Pro Leu Met Ala Ile Val Leu Glu Ile Ala Met 10 5 Ile Ala Ser Phe Gly Leu Phe Val Glu Tyr Glu Thr Asp His Thr Val 20 25 Leu Glu His Phe Asn Ile Thr Lys Pro Ser Asp Met Gly Ile Phe Phe 40 45 Glu Leu Tyr Pro Leu Phe Gln Asp Val His Gly Met Ile Phe Val Gly 55 60 Phe Asp Phe Pro Pro Asp Leu Pro Glu Glu Leu Trp Val Ser Gln Arg 70 75 Gly Tyr * 82

<210> 1717 <211> 57 <212> PRT <213> Homo sapiens

 $<\!400>$ 1717 Met Ala Leu Phe Phe Leu Ala Leu Asn Phe Trp Lys Val Gly Met Ala

<210> 1718 <211> 76 <212> PRT <213> Homo sapiens

<210> 1719 <211> 71 <212> PRT <213> Homo sapiens

<210> 1720 <211> 101 <212> PRT <213> Homo sapiens

 Phe
 Pro
 Leu
 Pro
 His
 Pro
 Thr
 Leu
 Gly
 Pro
 Arg
 Arg
 His
 Ala
 Ser
 Leu

 Thr
 Gln
 Leu
 Gly
 Pro
 Ala
 Phe
 Trp
 Met
 Ala
 Trp
 Gly
 Arg
 Pro
 Trp
 Ala

 His
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 Gly
 Pro
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 Gly
 Gln
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 Gly
 Gln
 Leu
 Trp
 Leu
 Trp
 Leu
 Ser
 Ser
 Val

 Glu
 Glu
 His
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 Leu
 Ala
 Ala
 Trp
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 Gln
 Pro
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<210> 1721 <211> 48 <212> PRT <213> Homo sapiens

<210> 1722 <211> 70 <212> PRT <213> Homo sapiens

varione paparent

<210> 1723 <211> 54 <212> PRT <213> Homo sapiens

 $<\!400>$ 1723 Met Asp Leu Ile Phe Val Lys Val Leu Leu Ile Phe Ala Ala Ile Gln

<210> 1724 <211> 60 <212> PRT <213> Homo sapiens

Leu Asn His Ile Met Leu Thr Thr Lys Phe Trp
50 55 59

<210> 1725 <211> 63 <212> PRT <213> Homo sapiens

<210> 1726 <211> 57 <212> PRT <213> Homo sapiens

Ser Gln Arg Leu Lys Glu Glu * 50 55 56

<210> 1727

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1727

 Met Arg Trp Pro Trp Ala Ser Trp Ala Ala Val Leu Leu Lys Leu Pro

 1
 5
 10
 15

 Arg Arg Val Leu Pro Trp Leu Pro Cys Gly His Gln Gln His Val Arg
 20
 25
 30

 Ala Thr Ala Ser Ser Arg Ser Pro Pro Met Pro Val Thr Lys
 45
 46

<210> 1728

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1728

<210> 1729

<211> 49

<212> PRT

<213> Homo sapiens

<400> 1729

 Met
 Val
 Leu
 Leu
 Pro
 Leu
 Gln
 Cys
 Gly
 Leu
 Thr
 Lys
 Ala
 Ser
 Ser
 Cys

 Leu
 His
 Thr
 Leu
 Cys
 Ser
 Ser
 Ser
 Asp
 Gln
 Ile
 Gly
 Tyr
 Leu
 Pro
 Val

 Lys
 Asn
 Thr
 Asp
 Gln
 Leu
 Gly
 Leu
 Gln
 Met
 Glu
 Val
 Ala
 Glu
 Met
 Cys

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 48
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<210> 1730

<211> 50

<212> PRT

<213> Homo sapiens

<210> 1731 <211> 227 <212> PRT <213> Homo sapiens

<400> 1731 Met Gly Cys Asp Gly Arg Val Ser Gly Leu Leu Arg Arg Asn Leu Gln 10 Pro Thr Leu Thr Tyr Trp Ser Val Phe Phe Ser Phe Gly Leu Cys Ile 25 Ala Phe Leu Gly Pro Thr Leu Leu Asp Leu Arg Cys Gln Thr His Ser 40 Ser Leu Pro Gln Ile Ser Trp Val Phe Phe Ser Gln Gln Leu Cys Leu . 55 60 Leu Leu Gly Ser Ala Leu Gly Gly Val Phe Lys Arg Thr Leu Ala Gln 75 Ser Leu Trp Ala Leu Phe Thr Ser Ser Leu Ala Ile Ser Leu Val Phe 90 Ala Val Ile Pro Phe Cys Arg Asp Val Lys Val Leu Ala Ser Val Met 105 Ala Leu Ala Gly Leu Ala Met Gly Cys Ile Asp Thr Val Ala Asn Met 120 Gln Leu Val Arg Met Tyr Gln Lys Asp Ser Ala Val Phe Leu Gln Val 135 Leu His Phe Phe Val Gly Phe Gly Ala Leu Leu Ser Pro Leu Ile Ala 150 155 Asp Pro Phe Leu Ser Glu Ala Asn Cys Leu Pro Ala Asn Ser Thr Gly 165 170 Gln His His Leu Pro Arg Ala Thr Cys Ser Met Ser Pro Gly Cys Trp 185 Gly Gln His His Val Asp Ala Gln Ala Leu Val Gln Pro Asp Val Pro 200 Lys Ala Asp Ser Gln Gly Pro Gly Arg Glu Pro Glu Gly Pro Met Pro 215 Ser Gly * 225 226

<210> 1732 <211> 102 <212> PRT <213> Homo sapiens

<210> 1733 <211> 139 <212> PRT <213> Homo sapiens

<400> 1733 Met Lys Phe Thr Thr Leu Leu Phe Leu Ala Ala Val Ala Gly Ala Leu 10 Val Tyr Ala Glu Asp Ala Ser Ser Asp Ser Thr Gly Ala Asp Pro Ala 25 Gln Glu Ala Gly Thr Ser Lys Pro Asn Glu Glu Ile Ser Gly Pro Ala 40 Glu Pro Ala Ser Pro Pro Glu Thr Thr Thr Ala Gln Glu Thr Ser 55 Ala Ala Val Gln Gly Thr Ala Lys Val Thr Ser Ser Arg Gln Glu 70 75 Leu Asn Pro Leu Lys Ser Ile Val Glu Lys Ser Ile Leu Leu Thr Glu 90 85 Gln Ala Leu Ala Lys Ala Gly Lys Gly Met His Gly Gly Val Pro Gly 105 Gly Lys Gln Phe Ile Glu Asn Gly Ser Glu Phe Ala Gln Lys Leu Leu 120 Lys Lys Phe Ser Leu Leu Lys Pro Trp Ala 130 135

<210> 1734 <211> 60 <212> PRT <213> Homo sapiens

35 40 45
Gln Leu Val Cys Trp Ile Leu Thr Phe Phe *
50 55 59

<210> 1735 <211> 73 <212> PRT

<213> Homo sapiens

<400> 1735

 Met
 Cys
 Ala
 Cys
 Ala
 Val
 Arg
 Ala
 Leu
 Ser
 Leu
 Ala
 Cys
 Ala
 Cys
 Ala
 Arg
 Ala
 Arg
 Ala
 Pro
 Arg
 Arg
 Tyr
 Val
 Gly

 Gly
 Glu
 Arg
 Arg
 Val
 Gln
 Ser
 Pro
 Ala
 Arg
 Pro
 Ala
 Asp
 Ser
 Val
 Ala

 Arg
 Ile
 Ala
 Phe
 Arg
 Phe
 Arg
 Thr
 Asp
 Leu
 Gly
 Ser
 Gly

 Pro
 Ser
 Leu
 His
 Leu
 Gly
 Ile
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<210> 1736 <211> 65 <212> PRT <213> Homo sapiens

<210> 1737 <211> 47 <212> PRT <213> Homo sapiens

<210> 1738 <211> 107 <212> PRT <213> Homo sapiens

<400> 1738 Met Val Thr Gln Leu Thr Leu Glu Val Leu His Leu Ser Leu Val Val 5 7.0 Gly Gln Val Ser Asn Asn Leu Leu His Ile Gly Pro Leu Ala Ser 2.0 25 Glu Gln Met Phe Tyr Ala Val Ala Thr Lys Ile Arg Asp Glu Asn Thr 35 40 Tyr Lys Ile Cys Thr Trp Leu Glu Ile Lys Val His His Val Leu Leu 55 60 His Ile Gln Gly Thr Leu Thr Cys Ser Tyr Leu Ser His Ser Glu Gln 70 75 Leu Val Phe Gln Ser Tyr Glu Tyr Val Asp Cys Arg Gly Asn Ala Ser 90 Val Pro His Gln Leu Thr Pro His Pro Pro 105 106

<210> 1739 <211> 90 <212> PRT <213> Homo sapiens

<210> 1740 <211> 57 <212> PRT <213> Homo sapiens

 $<\!400>$ 1740 Met His Cys Val Leu Glu Ile Leu Val Ser Val Leu Gly Leu Thr His 1 5 10 15 His Leu Leu Leu Arg Asp Arg Asp His Tyr Arg Leu Val Arg Leu Met

20 25 30

Gly Asp Val Gly Glu Gly Glu Leu Lys Ala Met Trp Arg Val Cys
35 40 45

Leu Ser Val Cys Arg Val Asp Lys *
50 55 56

<210> 1741 <211> 49 <212> PRT <213> Homo sapiens

<210> 1742 <211> 87 <212> PRT <213> Homo sapiens

<400> 1742 Met Ser Phe Val Lys Ile Leu Ile Trp Glu Leu Phe Ile Ala Cys Phe 1 5 10 Pro Gln Gly Pro Leu Val His Ser Gly Lys Met Leu Lys His Gly Leu 20 25 Asp Trp His Arg Thr Leu Leu Gln Lys His Pro Cys Ile Leu Phe Phe 40 Ser Phe Leu Lys Trp Asn Leu Ala Leu Ser Pro Trp Met Glu Gly Ser 60 55 Gly Ala Ile Ser Ala His Cys Asn Leu Cys Leu Leu Gly Ser Arg Asp 70 Ala Pro Ala Ser Val Ser 85 86

<210> 1743 <211> 49 <212> PRT <213> Homo sapiens

<400> 1743

Met Gly Phe Leu Ser Leu Thr Leu Tyr Leu Leu Thr Ser Leu Asn Lys
1 5 10 15

Met Leu Phe Lys Leu Arg Gly Ala Gln Pro Thr Glu Glu Asp Ile Gly
20 25 30

Gly Trp Leu Asn Glu Leu Lys Thr Ser Leu Lys Tyr Ile Arg Leu Arg
35 40 45 48

<210> 1744

<211> 57

<212> PRT

<213> Homo sapiens

<400> 1744

<210> 1745

<211> 96

<212> PRT

<213> Homo sapiens

<400> 1745

 Met
 Asn
 Gln
 Leu
 Ser
 Phe
 Leu
 Leu
 Phe
 Leu
 Ile
 Ala
 Thr
 Thr
 Arg
 Gly

 Trp
 Ser
 Thr
 Asp
 Glu
 Ala
 Asn
 Thr
 Tyr
 Phe
 Leu
 Glu
 Cys
 Thr
 Cys
 Ser

 Trp
 Ser
 Pro
 Ser
 Leu
 Pro
 Lys
 Ser
 Cys
 Pro
 Glu
 Ile
 Lys
 Asp
 Gln
 Cys

 Pro
 Ser
 Ala
 Phe
 Asp
 Gly
 Leu
 Tyr
 Phe
 Ile
 Arg
 Thr
 Glu
 Asn
 Ala
 Val
 Val
 Val
 Fro
 Fro
 Ile
 Arg
 Thr
 Glu
 Asn
 Ala
 Val
 Val
 Val
 Fro
 Fro
 Fro
 Ile
 Arg
 Thr
 Glu
 Asn
 Ala
 Val
 Arg
 Ile
 Arg
 Thr
 Thr
 Thr
 Thr

<210> 1746

<211> 53

<212> PRT

<213> Homo sapiens

<400> 1746

Met Val Ile Ser Ala Ala Val Leu Ser Ser Ile Leu Cys Val Phe Leu 1 5 5 5 10 10 5 15 Ser Lys Leu Val Leu Met Asn Asp Glu Cys Leu Arg Leu Thr Phe Trp 20 5 25 5 30 Leu His Cys Asn Ala Lys His Tyr Arg Tyr Ser Met Leu Gly Phe Pro

35 40 45 Lys Leu Thr Ser Val 50 53

<210> 1747 <211> 49 <212> PRT <213> Homo sapiens

<210> 1748 <211> 196 <212> PRT <213> Homo sapiens

apromo papromo

<400> 1748 Met Ala Met Leu Pro Phe Pro Ile Phe Leu Val Leu Leu Arg Gly 1.0 Leu Val Leu Trp Thr Pro Ala Ser Ser Gly Thr Ile Met Pro Glu Glu 25 Arg Lys Thr Glu Ile Glu Arg Glu Thr Glu Thr Glu Ser Glu Thr Val 40 Ile Gly Thr Glu Lys Glu Asn Ala Pro Glu Arg Glu Arg Gly Ser Val 55 Ile Thr Val Leu His Gln Val Phe Ser Thr Ala Met Lys Asn Asp Thr 75 70 Asp Thr Gly Asn Met Gln Lys Glu Val Met Ser Val Thr Glu Gln Val 90 Glu Lys Lys Lys Asn Asp Ile Glu Lys Asp Asp Thr Gly Arg Lys Arg 100 105 Lys Pro Asp Ile Ser Leu Leu Glu Val Ile Val Asp Val Ala Met Lys 120 115 Val Lys Lys Glu Ile Val Thr Gly Asp Thr Asn Thr Lys Asn Leu Lys 135 140 Glu Ala Lys Lys Glu Lys Lys Arg Ala Val Ser Leu Pro Leu Asn Arg 155 Arg Ala Pro Lys Leu His Leu Gln Asn Arg His Gly Phe Gly Leu Leu 170 Cys Ile Leu Val Pro Glu Val Asp Thr Ile Asn Leu Val Ile Phe Leu 185 Asp Asn Val 195

<210> 1749 <211> 46 <212> PRT <213> Homo sapiens

<210> 1750 <211> 82 <212> PRT <213> Homo sapiens

<210> 1751 <211> 94 <212> PRT <213> Homo sapiens

<400> 1751 Met Gly Ser Val Phe Trp His Val Leu Phe Cys Ile Ser Gly Val Cys 5 10 Leu Trp Cys Ala His Arg Met Ala Ala Phe Leu Gln Gln Met Ala Val 25 20 Leu Leu Pro Val Asp Cys Glu Arg Pro Ala Ala Val His Trp Leu Ala 40 Leu Cys Gly Cys Cys Tyr Gly Gln Leu Val Trp Glu Ser Arg Thr Arg 55 Ser Cys Phe Trp Ser Leu Glu Cys Leu Cys Phe Gly Gln His Phe 75 70 Gly Ser Val Pro Ser Phe Phe Cys Ser Ser Val Trp Leu * 90 85

<210> 1752 <211> 143 <212> PRT <213> Homo sapiens

<400> 1752 Met Asp Thr Trp Leu Val Cys Trp Ala Ile Phe Ser Leu Leu Lys Ala 10 Gly Leu Thr Glu Pro Glu Val Thr Gln Thr Pro Ser His Gln Val Thr 25 Gln Met Gly Gln Glu Val Ile Leu Arg Cys Val Pro Ile Ser Asn His 40 Leu Tyr Phe Tyr Trp Tyr Arg Gln Ile Leu Gly Gln Lys Val Glu Phe 55 Leu Val Ser Phe Tyr Asn Asn Glu Ile Ser Glu Lys Ser Glu Ile Phe · 70 75 Asp Asp Gln Phe Ser Val Glu Arg Pro Asp Gly Ser Asn Phe Thr Leu 85 90 Lys Ile Arg Ser Thr Lys Leu Glu Asp Ser Ala Met Tyr Phe Cys Ala 100 105 Ser Ser Glu Arg Gly Ser Gly Ala Asn Val Leu Thr Phe Gly Ala Gly 120 125 Ser Arg Leu Thr Val Leu Glu Asp Leu Lys Asn Val Phe Pro Pro 130 1.35 140

<210> 1753 <211> 64 <212> PRT <213> Homo sapiens

<210> 1754 <211> 124 <212> PRT <213> Homo sapiens

 $<\!400\!>$ 1754 Met Val Leu Gln Thr His Ala Phe Ile Ser Leu Leu Leu Trp Ile Ser 1 5 10 15 Gly Ala Cys Gly Asp Ile Val Met Thr His Ser Pro Asp Ser Leu Ala 20 25 30

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 Val
 Ser
 Leu
 Gly
 Glu
 Thr
 Ala
 Thr
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 Asp
 Cys
 Arg
 Ser
 Ser
 Gln
 Ser

 Val
 Leu
 Tyr
 His
 Ala
 Asn
 Asn
 Lys
 Asn
 Tyr
 Leu
 Thr
 Trp
 Tyr
 Gln
 Gln
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 Arg
 Pro
 Arg
 Pro
 Lys
 Val
 Leu
 Ile
 Phe
 Trp
 Ala
 Ser
 Thr
 Arg

 65
 70
 75
 80

 Glu
 Thr
 Gly
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 Asp
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 Phe
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 Gly
 Ser
 Gly
 Thr
 Asp

 Glu
 Thr
 Gly
 Val
 Pro
 Asp
 Arg
 Phe
 Thr
 Gly
 Ser
 Gly
 Thr
 Asp
 90
 95
 90
 110
 110

<210> 1755 <211> 111 <212> PRT

<213> Homo sapiens

<400> 1755 Met Gln Ala Thr Ser Asn Leu Leu Asn Leu Leu Leu Leu Ser Leu Phe 10 Ala Gly Leu Asn Pro Ser Lys Thr His Ile Asn Pro Lys Glu Gly Trp 25 Gln Val Tyr Ser Ser Ala Gln Asp Pro Asp Gly Arg Gly Ile Cys Thr 40 Val Val Ala Pro Glu Gln Asn Leu Cys Ser Arg Asp Ala Lys Ser Arg 55 Gln Leu Arg Gln Leu Leu Glu Lys Val Gln Asn Met Ser Gln Ser Ile 75 Glu Val Leu Asn Leu Arg Thr Gln Arg Asp Phe Gln Tyr Val Leu Lys 90 Met Glu Thr Gln Met Lys Gly Leu Lys Ala Lys Phe Arg Gln Ile 105 1.00

<210> 1756 <211> 74 <212> PRT <213> Homo sapiens

<210> 1757 <211> 50 <212> PRT <213> Homo sapiens

<210> 1758 <211> 123 <212> PRT <213> Homo sapiens

<400> 1758 Met Ala Trp Ile Pro Leu Phe Leu Gly Val Leu Ala Tyr Cys Thr Glu 10 1 5 Ser Val Ala Ser Tyr Glu Leu Phe Gln Pro Pro Ser Val Ser Val Ser 25 Pro Gly Gln Thr Ala Thr Phe Thr Cys Ser Gly Asp Asp Leu Gly Asn 40 45 Lys Tyr Ile Cys Trp Tyr Leu Gln Lys Pro Gly Gln Pro Pro Val Val 55 60 Leu Met Tyr Gln Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe 70 75 Ser Gly Ser Asn Ser Gly Ser Thr Ala Thr Leu Thr Ile Ser Gly Thr 90 Gln Ala Thr Asp Glu Ala Leu Tyr Phe Cys Gln Ala Trp Asp Thr Asn 105 Gly Ala Val Phe Gly Gly Gly Thr Gln Leu Thr

<210> 1759 <211> 75 <212> PRT <213> Homo sapiens

Pro Cys Leu Tyr Leu Glu Gly Asn Pro Thr * 65 70 74

<210> 1760 <211> 122 <212> PRT <213> Homo sapiens

<400> 1760 Met Arg Leu Pro Asp Val Gln Leu Trp Leu Val Leu Leu Trp Ala Leu 10 Val Arg Ala Gln Gly Thr Gly Ser Val Cys Pro Ser Cys Gly Gly Ser 25 20 Lys Leu Ala Pro Gln Ala Glu Arg Ala Leu Val Leu Glu Leu Ala Lys 40 45 Gln Gln Ile Leu Asp Gly Leu His Leu Thr Ser Arg Pro Arg Ile Thr 55 60 His Pro Pro Pro Gln Ala Ala Leu Thr Arg Ala Leu Arg Arg Leu Gln 70 75 Pro Gly Ser Val Ala Pro Gly Asn Gly Glu Glu Val Ile Ser Phe Ala 85 90 Thr Val Thr Asp Ser Thr Ser Ala Tyr Ser Ser Leu Leu Thr Phe His 105 Leu Ser Thr Pro Arg Ser His His Leu Tyr

<210> 1761 <211> 123 <212> PRT <213> Homo sapiens

<400> 1761 Met Arg Val Arg Ile Gly Leu Thr Leu Leu Cys Ala Val Leu Leu 10 Ser Leu Ala Ser Ala Ser Ser Asp Glu Glu Gly Ser Gln Asp Glu Ser 20 25 Leu Asp Ser Lys Thr Thr Leu Thr Ser Asp Glu Ser Val Lys Asp His 40 Thr Thr Ala Gly Arg Val Val Ala Gly Gln Ile Phe Leu Asp Ser Glu 55 60 Glu Ser Glu Leu Glu Ser Ser Ile Gln Glu Glu Glu Asp Ser Leu Lys Ser Gln Glu Gly Glu Ser Val Thr Glu Asp Ile Ser Phe Leu Glu Ser 90 Pro Asn Pro Glu Asn Lys Asp Tyr Glu Glu Pro Lys Lys Val Arg Lys 105 Pro Gly Ser Leu Asp Ile Phe Leu Ala Phe *.. 120 122

<210> 1762 <211> 145

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<212> PRT
<213> Homo sapiens
<221> misc_feature
<222> (1)...(145)
<223> Xaa = any amino acid or nothing
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<400> 1762 Met Ala Leu Ala Leu Met Ile Ala Leu Gly Ser Leu Gly Leu His 10 Thr Trp Gln Ala Gln Ala Val Pro Thr Ile Leu Pro Leu Gly Leu Ala 20 25 Pro Asp Thr Phe Asp Asp Thr Tyr Val Gly Cys Ala Glu Glu Met Glu 40 Glu Lys Ala Ala Pro Leu Leu Lys Glu Glu Met Ala His His Ala Leu 55 Leu Arg Glu Ser Trp Glu Ala Ala Gln Glu Thr Trp Glu Asp Lys Arg 75 Arg Gly Leu Thr Leu Pro Pro Gly Phe Lys Ala Gln Asn Gly Ile Ala 85 Ile Met Val Tyr Thr Asn Ser Ser Asn Thr Leu Tyr Trp Glu Leu Asn 100 105 Xaa Ala Val Arg Thr Gly Gly Gly Ser Arg Glu Leu Tyr Met Arg His 120 125 Phe Pro Phe Lys Ala Leu His Phe Tyr Leu Ile Arg Ala Leu Gln Leu 135 Leu 145

<210> 1763 <211> 257 <212> PRT <213> Homo sapiens

<400> 1763 Met Lys Arg Glu Arg Gly Ala Leu Ser Arg Ala Ser Arg Ala Leu Arg 10 Leu Ala Pro Phe Val Tyr Leu Leu Ile Gln Thr Asp Pro Leu Glu 25 Gly Val Asn Ile Thr Ser Pro Val Arg Leu Ile His Gly Thr Val Gly 40 Lys Ser Ala Leu Leu Ser Val Gln Tyr Ser Ser Thr Ser Ser Asp Arg 55 Pro Val Val Lys Trp Gln Leu Lys Arg Asp Lys Pro Val Thr Val Val 75 Gln Ser Ile Gly Thr Glu Val Ile Gly Thr Leu Arg Pro Asp Tyr Arg 90 Asp Arg Ile Arg Leu Phe Glu Asn Gly Ser Leu Leu Leu Ser Asp Leu 105 Gln Leu Ala Asp Glu Gly Thr Tyr Glu Val Glu Ile Ser Ile Thr Asp 120 Asp Thr Phe Thr Gly Glu Lys Thr Ile Asn Leu Thr Val Asp Val Pro 135 140 Ile Ser Arg Pro Gln Val Leu Gly Ala Ser Thr Thr Val Leu Glu Leu 150 155

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Ser Glu Ala Phe Thr Leu Asn Cys Ser His Glu Asn Gly Thr Lys Pro
                165
                                    170
 Ser Tyr Thr Trp Leu Lys Asp Gly Lys Pro Leu Leu Asn Asp Ser Arg
                                 185
 Met Leu Leu Ser Pro Asp Gln Lys Val Leu Thr Ile Thr Arg Val Leu
                                                 205
                             200
 Met Glu Asp Asp Asp Leu Tyr Ser Cys Val Val Glu Asn Pro Ile Asn
                         215
                                             220
 Gln Gly Arg Thr Leu Pro Cys Lys Ile Thr Glu Tyr Arg Lys Ser Ser
                    230
                                        235
 Leu Ser Ser Ile Trp Leu Gln Glu Ala Phe Ser Ser Leu Gly Pro Trp
                 245
                                     250
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<210> 1764 <211> 166 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(166) <223> Xaa = any amino acid or nothing

<400> 1764 Met Ala Leu Lys Val Leu Leu Glu Glu Glu Lys Thr Phe Phe Thr Leu 10 Leu Val Leu Leu Gly Tyr Leu Ser Cys Lys Val Thr Cys Glu Ser Gly 20 25 Asp Cys Arg Gln Glu Phe Arg Asp Arg Ser Gly Asn Cys Val Pro 40 Cys Asn Gln Cys Gly Pro Gly Met Glu Leu Ser Lys Glu Cys Gly Phe 55 Gly Tyr Gly Glu Asp Ala Gln Cys Val Thr Cys Arg Leu His Arg Phe 70 75 Lys Glu Asp Trp Gly Phe Gln Lys Cys Lys Pro Cys Leu Asp Cys Ala 85 90 Val Val Asn Arg Phe Gln Lys Ala Asn Cys Ser Ala Thr Ser Asp Ala 100 105 Ile Cys Gly Asp Cys Leu Pro Gly Phe Tyr Arg Lys Thr Lys Leu Val 120 Gly Phe Gln Asp Met Glu Trp Trp Xaa Ala Leu Val Gly Arg Thr Pro 135 Phe Leu Pro Ser Leu Tyr Gly Asn Pro Ala Leu Gly Cys Gln Pro Arg 150 Val Gln Thr Phe Gly Glu 165 166

<210> 1765 <211> 90 <212> PRT <213> Homo sapiens

<210> 1766 <211> 57 <212> PRT <213> Homo sapiens

<210> 1767 <211> 63 <212> PRT <213> Homo sapiens

<210> 1768 <211> 174 <212> PRT <213> Homo sapiens

<400> 1768

Met Pro Ser Gly Cys Arg Cys Leu His Leu Val Cys Leu Leu Cys Ile 10 Leu Gly Ala Pro Gly Gln Pro Val Arg Ala Asp Asp Cys Ser Ser His Cys Asp Leu Ala His Gly Cys Cys Ala Pro Asp Gly Ser Cys Arg Cys Asp Pro Gly Trp Glu Gly Leu His Cys Glu Arg Cys Val Arg Met Pro Gly Cys Gln His Gly Thr Cys His Gln Pro Trp Gln Cys Ile Cys His 70 Ser Gly Trp Ala Gly Lys Phe Cys Asp Lys Asp Glu His Ile Cys Thr 85 90 Thr Gln Ser Pro Cys Gln Asn Gly Gly Gln Cys Met Tyr Asp Gly Gly 100 105 Gly Glu Tyr His Cys Val Cys Leu Pro Gly Phe His Gly Arg Asp Cys 120 125 Glu Arg Lys Ala Gly Pro Cys Glu Gln Ala Gly Ser Pro Cys Arg Asn 135 140 Gly Gly Gln Cys Gln Asp Asp Gln Gly Phe Ala Leu Asn Phe Thr Cys 150 155 Arg Cys Leu Val Gly Phe Val Gly Ala Arg Cys Asp Val *

<210> 1769 <211> 78

<212> PRT

<213> Homo sapiens

<400> 1769

<210> 1770

<211> 149

<212> PRT

<213> Homo sapiens

<400> 1770

 Met Leu Val Thr
 Leu Gly Leu Leu Thr
 Ser Phe Phe Ser Phe Leu Tyr

 1
 5
 10
 15

 Met Val Ala Pro Ser Ile Arg Lys Phe Phe Ala Gly Gly Val Cys Arg
 20
 25
 30

 Thr Asn Val Gln Leu Pro Gly Lys Val Val Val Ile Thr Gly Ala Asn
 35
 40
 45

 Thr Gly Ile Gly Lys Glu Thr Ala Arg Glu Leu Ala Ser Arg Gly Ala

55 Arg Val Tyr Ile Ala Cys Arg Asp Val Leu Lys Gly Glu Ser Ala Ala 75 70 Ser Glu Ile Arg Val Asp Thr Lys Asn Ser Gln Val Leu Val Arg Lys 85 90 Leu Asp Leu Ser Asp Thr Lys Ser Ile Arg Ala Phe Ala Glu Gly Phe 105 100 Leu Ala Glu Glu Lys Gln Leu His Ile Leu Ile Asn Asn Ala Gly Val 120 125 Met Met Cys Pro Tyr Ser Lys Thr Ala Asp Gly Phe Glu Thr His Leu 135 140 Gly Val Asn His Leu 149

<210> 1771 <211> 76 <212> PRT <213> Homo sapiens

<210> 1772 <211> 128 <212> PRT <213> Homo sapiens

<210> 1773 <211> 614 <212> PRT <213> Homo sapiens

<400> 1773 Met Gly Ala Leu Arg Pro Thr Leu Leu Pro Pro Ser Leu Pro Leu Leu 10 Leu Leu Met Leu Gly Met Gly Cys Trp Ala Arg Glu Val Leu Val 20 25 Pro Glu Gly Pro Leu Tyr Arg Val Ala Gly Thr Ala Val Ser Ile Ser 40 Cys Asn Val Thr Gly Tyr Glu Gly Pro Ala Gln Gln Asn Phe Glu Trp Phe Leu Tyr Arg Pro Glu Ala Pro Asp Thr Ala Leu Gly Ile Val Ser 70 75 Thr Lys Asp Thr Gln Phe Ser Tyr Ala Val Phe Lys Ser Arg Val Val 85 90 Ala Gly Glu Val Gln Val Gln Arg Leu Gln Gly Asp Ala Val Leu 105 Lys Ile Ala Arg Leu Gln Ala Gln Asp Ala Gly Ile Tyr Glu Cys His 120 Thr Pro Ser Thr Asp Thr Arg Tyr Leu Gly Ser Tyr Ser Gly Lys Val 135 140 Glu Leu Arg Val Leu Pro Asp Val Leu Gln Val Ser Ala Ala Pro Pro 150 155 Gly Pro Arg Gly Arg Gln Ala Pro Thr Ser Pro Pro Arg Met Thr Val 165 170 His Glu Gly Gln Glu Leu Ala Leu Gly Cys Leu Ala Arg Thr Ser Thr 185 Gln Lys His Thr His Leu Ala Val Ser Phe Gly Arg Ser Val Pro Glu 200 Ala Pro Val Gly Arg Ser Thr Leu Gln Glu Val Val Gly Ile Arg Ser 215 Asp Leu Ala Val Glu Ala Gly Ala Pro Tyr Ala Glu Arg Leu Ala Ala Gly Glu Leu Arg Leu Gly Lys Glu Gly Thr Asp Arg Tyr Arg Met Val 245 250 Val Gly Gly Ala Gln Ala Gly Asp Ala Gly Thr Tyr His Cys Thr Ala 265 270 Ala Glu Trp Ile Gln Asp Pro Asp Gly Ser Trp Ala Gln Ile Ala Glu 280 285 Lys Arg Ala Val Leu Ala His Val Asp Val Gln Thr Leu Ser Ser Gln 295 300 Leu Ala Val Thr Val Gly Pro Gly Glu Arg Arg Ile Gly Pro Gly Glu 310 315 Pro Leu Glu Leu Leu Cys Asn Val Ser Gly Ala Leu Pro Pro Ala Gly 325 330 Arg His Ala Ala Tyr Ser Val Gly Trp Glu Met Ala Pro Ala Gly Ala 345 Pro Gly Pro Gly Arg Leu Val Ala Gln Leu Asp Thr Glu Gly Val Gly 360 Ser Leu Gly Pro Gly Tyr Glu Gly Arg His Ile Ala Met Glu Lys Val

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375
                                         380
Ala Ser Arg Thr Tyr Arg Leu Arg Leu Glu Ala Ala Arg Pro Gly Asp
                                    395
                  390
Ala Gly Thr Tyr Arg Cys Leu Ala Lys Ala Tyr Val Arg Gly Ser Gly
                                 410
              405
Thr Arg Leu Arg Glu Ala Ala Ser Ala Arg Ser Arg Pro Leu Pro Val
                             425
His Val Arg Glu Glu Gly Val Val Leu Glu Ala Val Ala Trp Leu Ala
                         440
Gly Gly Thr Val Tyr Arg Gly Glu Thr Ala Ser Leu Leu Cys Asn Ile
                              460
                     455
Ser Val Arg Gly Gly Pro Pro Gly Leu Arg Leu Ala Ala Ser Trp Trp
                 470
                             475
Val Glu Arg Pro Glu Asp Gly Glu Leu Ser Ser Val Pro Ala Gln Leu
                                490
Val Gly Gly Val Gly Gln Asp Gly Val Ala Glu Leu Gly Val Arg Pro
                            505
Gly Gly Gly Pro Val Ser Val Glu Leu Val Gly Pro Arg Ser His Arg
                                            525
                         520
Leu Arg Leu His Ser Leu Gly Pro Glu Asp Glu Gly Val Tyr His Cys
                     535
Ala Pro Ser Ala Trp Val Gln His Ala Asp Tyr Ser Trp Tyr Gln Ala
                 550
                                    555
Gly Ser Ala Arg Ser Gly Pro Val Thr Val Tyr Pro Tyr Met His Ala
              565 . 570
Leu Asp Thr Leu Phe Val Pro Leu Leu Val Gly Thr Gly Val Ala Leu
                             585
Val Thr Gly Ala Thr Val Leu Gly Thr Ile Thr Cys Cys Phe Met Lys
                         600
       595
Arg Leu Arg Lys Arg *
    610
         613
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<210> 1774

<211> 156

<212> PRT

<213> Homo sapiens

<400> 1774 Met Glu Ala Leu Thr Leu Trp Leu Leu Pro Trp Ile Cys Gln Cys Val 5 10 Ser Val Arg Ala Asp Ser Ile Ile His Ile Gly Ala Ile Phe Glu Glu 25 20 Asn Ala Ala Lys Asp Asp Arg Val Phe Gln Leu Ala Val Ser Asp Leu 40 Ser Leu Asn Asp Asp Ile Leu Gln Ser Glu Lys Ile Thr Tyr Ser Ile 55 Lys Val Ile Glu Ala Asn Asn Pro Phe Gln Ala Val Gln Glu Ala Cys 75 70 Asp Leu Met Thr Gln Gly Ile Leu Ala Leu Val Thr Ser Thr Gly Cys 90 85 Ala Ser Ala Asn Ala Leu Gln Ser Leu Thr Asp Ala Met His Ile Pro 105 His Leu Phe Val Gln Arg Asn Pro Gly Gly Ser Pro Arg Thr Ala Cys 125 120 His Leu Asn Pro Ser Pro Asp Gly Glu Ala Tyr Thr Leu Ala Ser Arg 135

Pro Pro Val Arg Leu Asn Asp Val Met Leu Arg Leu 145 150 155 156

> <210> 1775 <211> 896 <212> PRT <213> Homo sapiens

<400> 1775 Met Gln Lys Ala Ser Val Leu Leu Phe Leu Ala Trp Val Cys Phe Leu Phe Tyr Ala Gly Ile Ala Leu Phe Thr Ser Gly Phe Leu Leu Thr Arg 2.0 25 Leu Glu Leu Thr Asn His Ser Ser Cys Gln Glu Pro Pro Gly Pro Gly 40 Ser Leu Pro Trp Gly Ser Gln Gly Lys Pro Gly Ala Cys Trp Met Ala 55 Ser Arg Phe Ser Arg Val Val Leu Val Leu Ile Asp Ala Leu Arg Phe 70 75 Asp Phe Ala Gln Pro Gln His Ser His Val Pro Arg Glu Pro Pro Val 85 90 Ser Leu Pro Phe Leu Gly Lys Leu Ser Ser Leu Gln Arg Ile Leu Glu 105 Ile Gln Pro His His Ala Arg Leu Tyr Arg Ser Gln Val Asp Pro Pro 120 Thr Thr Met Gln Arg Leu Lys Ala Leu Thr Thr Gly Ser Leu Pro 135 Thr Phe Ile Asp Ala Gly Ser Asn Phe Ala Ser His Ala Ile Val Glu 150 155 Asp Asn Leu Ile Lys Gln Leu Thr Ser Ala Gly Arg Arg Val Val Phe 165 170 Met Gly Asp Asp Thr Trp Lys Asp Leu Phe Pro Gly Ala Phe Ser Lys 185 Ala Phe Phe Pro Ser Phe Asn Val Arg Asp Leu Asp Thr Val Asp 200 Asn Gly Ile Leu Glu His Leu Tyr Pro Thr Met Asp Ser Gly Glu Trp 215 220 Asp Val Leu Ile Ala His Phe Leu Gly Val Asp His Cys Gly His Lys 230 235 His Gly Pro His His Pro Glu Met Ala Lys Lys Leu Ser Gln Met Asp 245 250 Gln Val Ile Gln Gly Leu Val Glu Arg Leu Glu Asn Asp Thr Leu Leu 260 265 Val Val Ala Gly Asp His Gly Met Thr Thr Asn Gly Asp His Gly Gly 280 285 Asp Ser Glu Leu Glu Val Ser Ala Ala Leu Phe Leu Tyr Ser Pro Thr 295 300 Ala Val Phe Pro Ser Thr Pro Pro Glu Glu Pro Glu Val Ile Pro Gln 310 315 Val Ser Leu Val Pro Thr Leu Ala Leu Leu Gly Leu Pro Ile Pro 325 330 Phe Gly Asn Ile Gly Glu Val Met Ala Glu Leu Phe Ser Gly Gly Glu 345 Asp Ser Gln Pro His Ser Ser Ala Leu Ala Gln Ala Ser Ala Leu His 360 365 Leu Asn Ala Gln Gln Val Ser Arg Phe Phe His Thr Tyr Ser Ala Ala

	370					375					380				
		Asp	Leu	Gln	Ala 390		Glu	Leu	His	Gln 395		Gln	Asn	Leu	Phe 400
385 Ser	Lys	Ala	Ser			Tyr	Gln	Trp	Leu 410		Gln	Ser	Pro	Lys 415	
Ala	Glu	Ala	Thr	405 Leu	Pro	Thr	Val			Glu	Leu	Gln			Leu
Arg	Gly		420 Arg	Ala	Met	Cys		425 Glu	Ser	Trp	Ala		430 Phe	Ser	Leu
77-3	3	435	n 1 -	01	63	ml	440	T 033	7 011	ח ד ת	- ות	445	Caro	Dho	Tlo
	450		Ala	_	_	455					460				
Cys 465	Leu	Leu	Ala	Ser	Gln 470	Trp	Ala	Ile	Ser	Pro 475	Gly	Phe	Pro	Phe	Cys 480
Pro	Leu	Leu	Leu	Thr 485	Pro	Val	Ala	Trp	Gly 490	Leu	Val	Gly	Ala	Ile 495	Ala
Tyr	Ala	Gly	Leu 500		Gly	Thr	Ile	Glu 505		Lys	Leu	Asp	Leu 510	Val	Leu
Leu	Gly		Val	Ala	Ala	Val	Ser 520		Phe	Leu	Pro	Phe 525		Trp	Lys
Δla	Tro	515 Ala	Gly	Tro	Glv	Ser	-	Ara	Pro	Leu	Ala		Leu	Phe	Pro
1124	530				- 1	535	-2-	5			540				
Ile	Pro	Gly	Pro	Val	Leu	Leu	Leu	Leu	Leu	Phe	Arg	Leu	Ala	Val	Phe
545					550	_	_			555			_	_,	560
			Ser	565					570					575	
Leu	Gly	Ser	Phe 580	Ile	Leu	Leu	Leu	Val 585	Val	Gln	Leu	His	Trp 590	Glu	Gly
Gln	Leu	Leu 595	Pro	Pro	Lys	Leu	Leu 600	Thr	Met	Pro	Arg	Leu 605	Gly	Thr	Ser
Ala	Thr 610		Asn	Pro	Pro	Arg 615	His	Asn	Gly	Ala	Tyr 620	Ala	Leu	Arg	Leu
Gly 625		Gly	Leu	Leu	Leu 630		Thr	Arg	Leu	Ala 635	Gly	Leu	Phe	His	Arg 640
	Pro	Glu	Glu	Thr 645		Val	Cys	His	Ser 650		Pro	Trp	Leu	Ser 655	
Leu	Ala	Ser	Met		Gly	Gly	Arg	Ala 665		Asn	Leu	Trp	Tyr 670		Ala
Cys	Val		660 Ala	Leu	Val	Ala			Ala	Ala	Val			Trp	Leu
Arg	Arg	675 Tyr	Gly	Asn	Leu	Lys	680 Ser	Pro	Glu	Pro	Pro	685 Met	Leu	Phe	Val
71	690	C1	Leu	Dwo	T 011	695	ת 7 ת	Leu	G] w	Thr	700	בות	ጥኒም	Tro	Δla
705	ırp	СТУ	шеu	PIO	710	Mec	ALA	Бец	GLY	715	ALG	ALU	- y	1-6	720
	Ala	Ser	Gly	Ala 725		Glu	Ala	Pro	Pro 730	Arg	Leu	Arg	Val	Leu 735	Val
Ser	Gly	Ala	Ser 740		Val	Leu	Pro	Arg 745		Val	Ala	Gly	Leu 750	Ala	Ala
Ser	Gly	Leu 755	Ala	Leu	Leu	Leu	Trp 760		Pro	Val	Thr	Val 765		Val	Lys
Ala	_		Gly	Ala	Pro	Arg 775		Arg	Thr	Val	Leu 780		Pro	Phe	Ser
	770 Pro	Pro	Thr	Ser			Asp	Leu	Asp			Val	Pro	Gln	
785 Tvr	Δτα	Hie	Met	Gln	790 Glu	Glu	Phe	Arσ	G] v	795 Ara	Leu	Glu	Ara	Thr	800 Lys
- y -	T.A	****		805				9	810	=			3	815	4 -
Ser	Gln	Gly	Pro 820	Leu	Thr	Val	Ala	Ala 825	Tyr	Gln	Leu	Gly	Ser 830	Val	Tyr
Ser	Ala	Ala 835	Met	Val	Thr	Ala	Leu 840	Thr	Leu	Leu	Ala	Phe 845	Pro	Leu	Leu

<210> 1776 <211> 178 <212> PRT <213> Homo sapiens

<400> 1776 Met Trp Ala Cys Trp Cys Val Leu Gly Thr Pro Gly Val Ala Met Val Leu Leu His Thr Thr Ile Ser Phe Cys Val Ala Gln Phe Arg Ser Gln 25 Leu Leu Thr Trp Leu Cys Ser Leu Leu Leu Ser Thr Leu Arg Leu Gln Gly Val Glu Glu Val Lys Arg Arg Trp Tyr Lys Thr Glu Asn Glu 55 Tyr Tyr Leu Leu Gln Phe Thr Leu Thr Val Arg Cys Leu Tyr Tyr Thr 75 Ser Phe Ser Leu Glu Leu Cys Trp Gln Gln Leu Pro Ala Ala Ser Thr 90 Ser Tyr Ser Phe Pro Trp Met Leu Ala Tyr Val Phe Tyr Tyr Pro Val 105 110 Leu His Asn Gly Pro Ile Leu Ser Phe Ser Glu Phe Ile Lys Gln Arg 120 125 Ser Gln Trp Ser Asn Arg Glu Phe Gly Met Glu Val Glu Ser Lys Gly 135 140 Pro Gly Ala His Pro Pro Gly Phe Glu Ser Leu Leu Cys Phe Gly Leu 150 155 Arg Val Leu Ala Glu Leu Leu Thr Leu Leu Met Pro Gln Ser Ser Tyr 165 170 Gln * 177

<210> 1777 <211> 59 <212> PRT <213> Homo sapiens

PCT/US01/02687 WO 01/54477

50 55 59

<210> 1778 <211> 137 <212> PRT <213> Homo sapiens

<400> 1778 Met Val Ala Pro Gly Leu Val Leu Gly Leu Val Leu Pro Leu Ile Leu 10 Trp Ala Asp Arg Ser Ala Gly Ile Gly Phe Arg Phe Ala Ser Tyr Ile 20 25 Asn Asn Asp Met Val Leu Gln Lys Glu Pro Ala Gly Ala Val Ile Trp 40 Gly Phe Gly Thr Pro Gly Ala Thr Val Thr Val Thr Leu Arg Gln Gly 55 Gln Glu Thr Ile Met Lys Lys Val Thr Ser Val Lys Ala His Ser Asp 70 Thr Trp Met Val Val Leu Asp Pro Met Lys Pro Gly Gly Pro Phe Glu 90 Val Met Ala Gln Gln Thr Leu Glu Lys Ile Asn Phe Thr Leu Arg Val 105 His Asp Val Leu Phe Gly Asp Val Trp Leu Cys Ser Gly Gln Ser Asn 120 125 Met Gln Met Thr Val Leu Gln Ile Phe 135

<210> 1779 <211> 65 <212> PRT <213> Homo sapiens

<400> 1779

Met Lys Val Phe Phe Leu Asp Glu Ser Trp Pro Gln Trp Arg Phe Ala 1 5 10 Ala Gly Leu Leu Ala Leu Ser Phe Gly Gly Pro Ala Trp Lys Phe Leu 20 25 30 Ser Val Gln Arg Val Ile Pro Trp Leu Trp Ala Ala Lys Glu Lys Pro 40 Leu Gly Pro Leu Ala Thr Pro Pro Arg Leu Asn Pro Lys Val Gly Val 55 60

<210> 1780 <211> 53 <212> PRT <213> Homo sapiens

<400> 1780

<210> 1781 <211> 109 <212> PRT <213> Homo sapiens

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<400> 1781 Met Met His Asn Ile Ile Val Lys Glu Leu Ile Val Thr Phe Phe Leu 10 Gly Ile Thr Val Val Gln Met Leu Ile Ser Val Thr Gly Leu Lys Gly 20 25 Val Glu Ala Gln Asn Gly Ser Glu Ser Glu Val Phe Val Gly Lys Tyr 40 Glu Thr Leu Val Phe Tyr Trp Pro Ser Leu Leu Cys Leu Ala Phe Leu 55 60 Leu Gly Arg Phe Leu His Met Phe Val Lys Ala Leu Arg Val His Leu 70 75 Gly Trp Glu Leu Gln Val Glu Glu Lys Ser Val Leu Glu Val His Gln 90 Gly Glu His Val Lys Gln Leu Leu Arg Ile Pro Arg Pro 100 105

<210> 1782 <211> 58 <212> PRT <213> Homo sapiens

<210> 1783 <211> 102 <212> PRT <213> Homo sapiens

<400> 1783 Met Leu Ile Pro His Gln Leu Pro Leu Cys Ser Pro Trp Leu Val Gln 10 Ala Met Leu Thr Ile Glu Val Pro Trp Leu Leu Gly Leu Ala His Tyr 25 Arg Leu Gly Trp His Ala Leu Glu Gly Ile Phe Trp Trp Gly Ala Ser 40 Val Phe His Ala Leu Gln Ala Met Leu Val Arg Lys Trp Pro Leu Gly 55 Leu Val Glu Phe Thr Gly Thr Cys Gly Ile Leu Val Glu Val Ile Gly 70 75 Leu Trp Trp Gly Glu Gly Ser Thr Gly Asn Arg Trp Met Gly Leu Asn 85 90 Ser Thr Gly Gly Gln * 100 101

<210> 1784 <211> 243 <212> PRT <213> Homo sapiens

<400> 1784 Met Gly Glu Ala Ser Pro Pro Ala Pro Ala Arg Arg His Leu Leu Val 1 5 10 Leu Leu Leu Leu Ser Thr Leu Val Ile Pro Ser Ala Ala Pro 20 25 Ile His Asp Ala Asp Ala Gln Glu Ser Ser Leu Gly Leu Thr Gly Leu 40 Gln Ser Leu Leu Gln Gly Phe Ser Arg Leu Phe Leu Lys Gly Asn Leu 55 Leu Arg Gly Ile Asp Ser Leu Phe Ser Ala Pro Met Asp Phe Arg Gly 70 75 Leu Pro Gly Asn Tyr His Lys Glu Glu Asn Gln Glu His Gln Leu Gly 90 Asn Asn Thr Leu Ser Ser His Leu Gln Ile Asp Lys Met Thr Asp Asn 105 Lys Thr Gly Glu Val Leu Ile Ser Glu Asn Val Val Ala Ser Ile Gln 120 125 Pro Ala Glu Gly Ser Phe Glu Gly Asp Leu Lys Val Pro Arg Met Glu 135 Glu Lys Glu Ala Leu Val Pro Ile Gln Lys Ala Thr Asp Ser Phe His 150 155 Thr Glu Leu His Pro Arg Val Ala Phe Trp Ile Ile Lys Leu Pro Arg 165 170 Arg Arg Ser His Gln Asp Ala Leu Glu Gly Gly His Trp Leu Ser Glu 185 Lys Arg His Arg Leu Gln Ala Ile Arg Asp Gly Leu Arg Lys Gly Thr 200 205 His Lys Asp Val Leu Glu Glu Gly Thr Glu Ser Ser His Ser Arg 220 215 Leu Ser Pro Arg Lys Thr His Leu Leu Tyr Ile Leu Arg Pro Ser Arg 230 235 Gln Leu * 242

<210> 1785 <211> 158 <212> PRT <213> Homo sapiens

<400> 1785 Met Lys Ala Leu Leu Leu Val Leu Pro Trp Leu Ser Pro Ala Asn 1.0 Tyr Ile Asp Asn Val Gly Asn Leu His Phe Leu Tyr Ser Glu Leu Cys 25 20 Lys Gly Ala Ser His Tyr Gly Leu Thr Lys Asp Arg Lys Arg Arg Ser 40 Gln Asp Gly Cys Pro Asp Gly Cys Ala Ser Leu Thr Ala Thr Ala Pro 55 60 Ser Pro Glu Val Ser Ala Ala Ala Thr Ile Ser Leu Met Thr Asp Glu 75 Pro Gly Leu Asp Asn Pro Ala Tyr Val Ser Ser Ala Glu Asp Gly Gln 90 Pro Ala Ile Ser Pro Val Asp Ser Gly Arg Ser Asn Arg Thr Arg Ala 105 Arg Pro Phe Glu Arg Ser Thr Ile Ile Ser Arg Ser Phe Lys Lys Ile 120 Asn Arg Ala Leu Ser Val Leu Arg Arg Thr Lys Ser Gly Ser Ala Val 135 Ala Asn His Ala Asp Gln Gly Arg Glu Asn Ser Glu Asn Thr 155 150

<210> 1786 <211> 142 <212> PRT <213> Homo sapiens

<400> 1786 Met Glu Ser Ala Val Arg Val Glu Ser Gly Val Leu Val Gly Val Val 10 Cys Leu Leu Leu Ala Cys Pro Ala Thr Ala Thr Gly Pro Glu Val Ala 25 Gln Pro Glu Val Asp Thr Thr Leu Gly Arg Val Arg Gly Arg Gln Val Gly Val Lys Gly Thr Asp Arg Leu Val Asn Val Phe Leu Gly Ile Pro 55 Phe Ala Gln Pro Pro Leu Gly Pro Asp Arg Phe Ser Ala Pro His Pro 75 Ala Gln Pro Trp Glu Gly Val Arg Asp Ala Ser Thr Ala Pro Pro Met 90 85 Cys Leu Gln Asp Val Glu Ser Met Asn Ser Ser Arg Phe Val Leu Asn 105 100 Gly Lys Gln Gln Ile Phe Ser Val Ser Glu Asp Cys Leu Val Leu Asn 120 Val Tyr Ser Pro Ala Glu Val Pro Ala Gly Ser Gly Arg Pro 140 142 135 130

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<210> 1787
<211> 120
<212> PRT
<213> Homo sapiens

<221> misc_feature
<222> (1)...(120)
<223> Xaa = any amino acid or nothing
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<400> 1787 Met Ala Leu Thr Gly Tyr Ser Trp Leu Leu Ser Ala Thr Phe Leu 10 Asn Val Gly Ala Glu Ile Ser Ile Thr Leu Glu Pro Ala Gln Pro Ser 20 25 Glu Gly Asp Asn Val Thr Leu Val Val His Gly Leu Ser Gly Glu Leu 40 Leu Ala Tyr Ser Trp Tyr Ala Gly Pro Thr Leu Ser Val Ser Tyr Leu 55 60 Val Ala Ser Tyr Ile Val Ser Thr Gly Asp Glu Thr Pro Gly Pro Ala 70 75 His Thr Xaa Arg Glu Ala Val Arg Pro Asp Gly Ser Leu Asp Ile Gln 90 Gly Ile Leu Pro Arg His Ser Ser Thr Tyr Ile Leu Gln Thr Phe Asn 105 Arg Gln Leu Gln Thr Glu Val Gly

<210> 1788 <211> 68 <212> PRT <213> Homo sapiens

<210> 1789 <211> 133 <212> PRT <213> Homo sapiens

<400> 1789
Met Ala Val Val Ile Arg Leu Leu Gly Leu Pro Phe Ile Ala Gly Pro
1 5 10 15

Val Asp Ile Arg His Phe Phe Thr Gly Leu Thr Ile Pro Asp Gly Gly 25 Val His Ile Ile Gly Glu Ile Gly Glu Ala Phe Ile Ile Phe Ala Thr Asp Glu Asp Ala Arg Arg Ala Ile Ser Arg Ser Gly Gly Phe Ile 55 Lys Asp Ser Ser Val Glu Leu Phe Leu Ser Ser Lys Ala Glu Met Gln 70 Lys Thr Ile Glu Met Lys Arg Thr Asp Arg Val Gly Arg Gly Arg Pro Gly Ser Gly Thr Ser Gly Val Asp Ser Leu Ser Asn Phe Ile Glu Ser 100 105 Val Lys Glu Glu Ala Ser Asn Ser Gly Tyr Gly Ser Ser Ile Asn Gln 115 120 Asp Ala Gly Phe His 130 133

<210> 1790

<211> 82

<212> PRT

<213> Homo sapiens

<400> 1790

 Met
 Ala
 Ala
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 Cys
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<210> 1791

<211> 50

<212> PRT

<213> Homo sapiens

<400> 1791

 Met His Ala Ser Glu Gly Leu Pro Ala Leu Pro Leu Leu Ala Leu Val

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 10
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 Ser His Ser His Ser Cys Pro Pro Leu Pro Leu Gln Pro His His Leu
 20
 25
 30

 Pro Ala Ile Leu Phe Phe Leu Val Gly His Gln Leu Met Lys Cys Ile
 35
 40
 45

 Arg *
 49

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<210> 1792
<211> 166
<212> PRT
<213> Homo sapiens
<221> misc_feature
<222> (1)...(166)
<223> Xaa = any amino acid or nothing
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<400> 1792 Met Leu Leu Trp Leu Leu Leu Ile Leu Thr Pro Gly Arg Glu Gln 10 5 Ser Gly Val Ala Pro Lys Ala Val Leu Leu Leu Asp Pro Pro Trp Ser 25 Thr Ala Phe Lys Gly Glu Lys Val Ala Leu Ile Cys Ser Ser Ile Ser 40 His Ser Leu Ala Gln Gly Asp Thr Tyr Trp Tyr His Asp Glu Lys Leu 55 Leu Lys Ile Lys His Asp Lys Ile Gln Ile Thr Glu Pro Gly Asn Tyr 70 75 Gln Cys Lys Thr Arg Gly Ser Ser Leu Ser Asp Ala Val His Val Glu 85 90 Phe Ser Pro Asp Trp Leu Ile Leu Gln Ala Leu His Pro Val Phe Glu 105 Gly Asp Asn Val Ile Leu Arg Cys Gln Gly Lys Asp Asn Lys Asn Thr 120 125 His His Lys Val Tyr Tyr Lys Asp Gly Lys Gln Xaa Ser Asn Ser Tyr 135 140 Asn Leu Glu Lys Asn Thr Val Asp Ser Val Ser Arg Asp Asn Ser Pro Tyr Tyr Cys Ala Gly * 165

<210> 1793 <211> 146 <212> PRT <213> Homo sapiens

<400> 1793 Met Ala Thr Ala Ala Gln Gly Pro Leu Ser Leu Leu Trp Gly Trp Leu 1 5 10 Trp Ser Glu Arg Phe Trp Leu Pro Glu Asn Val Ser Trp Ala Asp Leu 25 20 Glu Gly Pro Ala Asp Gly Tyr Gly Tyr Pro Arg Gly Arg His Ile Leu 40 Ser Val Phe Pro Leu Ala Ala Gly Ile Phe Phe Val Arg Leu Leu Phe 60 Glu Arg Phe Ile Ala Lys Pro Cys Ala Leu Arg Ile Gly Ile Glu Asp 75 Ser Gly Pro Tyr Gln Ala Gln Pro Asn Ala Ile Leu Glu Lys Val Phe 90 Ile Ser Ile Thr Lys Tyr Pro Asp Lys Lys Arg Leu Glu Giy Leu Ser 105 Lys Gln Leu Asp Trp Asn Val Arg Lys Ile Gln Cys Trp Phe Arg His 120

Arg Arg Asn Gln Asp Lys Pro Pro Thr Leu Thr Lys Phe Cys Glu Ser 130 135 140

Met *

<210> 1794 <211> 151 <212> PRT <213> Homo sapiens

<400> 1794 Met Glu Arg Arg Leu Leu Gly Gly Met Ala Leu Leu Leu Gln 10 Ala Leu Pro Ser Pro Leu Ser Ala Arg Ala Glu Pro Pro Gln Asp Lys 20 25 Glu Ala Cys Val Gly Thr Asn Asn Gln Ser Tyr Ile Cys Asp Thr Gly 40 His Cys Cys Gly Gln Ser Gln Cys Cys Asn Tyr Tyr Tyr Glu Leu Trp 55 60 Trp Phe Trp Leu Val Trp Thr Ile Ile Ile Leu Ser Cys Cys 75 Val Cys His His Arg Arg Ala Lys His Arg Leu Gln Ala Gln Gln Arg 85 90 Gln His Glu Ile Asn Leu Ile Ala Tyr Arg Glu Ala His Asn Tyr Ser 105 Ala Leu Pro Phe Tyr Phe Arg Phe Leu Pro Asn Tyr Leu Leu Pro Pro 120 Tyr Glu Glu Val Val Asn Arg Pro Pro Thr Pro Pro Pro Pro Tyr Ser 135 Ala Phe Gln Leu Gln Gln Gln 150 151 145

<210> 1795 <211> 177 <212> PRT <213> Homo sapiens

<400> 1795

 Met
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100 105 110

Ala Leu Lys Ile Leu Met Leu Val Gly Val Gln Pro Ala Arg Leu Ile

PCT/US01/02687 WO 01/54477

120 115 Leu Gly Met Met Val Thr Thr Ser Phe Leu Ser Met Trp Leu Ser Asn . 135 140 Thr Ala Ser Thr Ala Met Met Leu Pro Ile Ala Asn Ala Ile Leu Lys 150 155 Ser Leu Phe Gly Gln Lys Glu Val Arg Lys Asp Pro Gln Pro Gly Glu 170 175 176 165

<210> 1796

<211> 98

<212> PRT

<213> Homo sapiens

<221> misc feature

<222> (1)...(98)

<223> Xaa = any amino acid or nothing

<400> 1796

Met His Pro Leu Pro Gly Tyr Trp Ser Cys Tyr Cys Leu Leu Leu 10 Phe Ser Leu Gly Val Gln Gly Ser Leu Gly Ala Pro Ser Ala Ala Pro 25 20 Glu Gln Val His Leu Ser Tyr Pro Gly Glu Pro Gly Ser Met Thr Val 40 Thr Trp Thr Thr Trp Val Pro Thr Arg Ser Glu Val Gln Phe Gly Leu 55 Gln Pro Ser Gly Pro Leu Pro Leu Arg Ala Gln Gly Thr Phe Val Pro 70 Phe Val Asp Xaa Gly Ile Leu Arg Arg Lys Leu Tyr Ile His Arg Val Thr Leu

98

<210> 1797

<211> 96

<212> PRT

<213> Homo sapiens

<400> 1797

Met Phe Leu Trp Leu Phe Leu Ile Leu Ser Ala Leu Ile Ser Ser Thr 5 10 Asn Ala Asp Ser Asp Ile Ser Val Glu Ile Cys Asn Val Cys Ser Cys 25 30 Val Ser Val Glu Asn Val Leu Tyr Val Asn Cys Glu Lys Val Ser Val 40 Tyr Arg Pro Asn Gln Leu Lys Pro Pro Trp Ser Asn Phe Tyr His Leu 55 Asn Phe Gln Asn Asn Phe Leu Asn Ile Leu Tyr Pro Asn Thr Phe Leu 70 75 Asn Phe Ser His Ala Val Ser Leu His Leu Gly Asn Asn Lys Leu Gln <210> 1798 <211> 91 <212> PRT <213> Homo sapiens

<400> 1798 Met Arg Pro Ala Leu Ala Val Gly Leu Val Phe Ala Gly Cys Cys Ser Asn Val Ile Phe Leu Glu Leu Leu Ala Arg Lys His Pro Gly Cys Gly 20 25 Asn Ile Val Thr Phe Ala Gln Phe Leu Phe Ile Ala Val Glu Gly Phe 35 40 Leu Phe Glu Ala Asp Leu Gly Arg Lys Pro Pro Ala Ile Pro Ile Arg . 60 55 Tyr Tyr Ala Ile Met Val Thr Met Phe Phe Thr Val Ser Val Val Asn 75 70 Asn Tyr Ala Leu Asn Leu Asn Ile Ala Met Pro 85

<210> 1799 <211> 77 <212> PRT <213> Homo sapiens

<400> 1799 Met Arg Ser Leu Val Trp Val Leu Ile Gln Gln Leu Thr Pro Leu Tyr 10 Lys Gly Glu Thr Trp Thr Gln Thr Cys Thr Glu Asp His Val Thr Met 20 25 Lys Ala Glu Ile Arg Val Met Leu Leu Glu Ala Arg Glu Asp Cys Gln 40 Leu Met Thr Lys Arg Ser Gln Glu Thr Gly Leu Gln Arg Ile Leu Pro 55 Glu Gly Ser Gln Lys Glu Pro Thr Leu Thr Thr Pro *

<210> 1800 <211> 182 <212> PRT

<213> Homo sapiens

<400> 1800 Met Ser Leu Lys Met Leu Ile Ser Arg Asn Lys Leu Ile Leu Leu Gly Ile Val Phe Phe Glu Arg Gly Lys Ser Ala Thr Leu Ser Leu Pro Lys Ala Pro Ser Cys Gly Gln Ser Leu Val Lys Val Gln Pro Trp Asn

40 Tyr Phe Asn Ile Phe Ser Arg Ile Leu Gly Gly Ser Gln Val Glu Lys 55 Gly Ser Tyr Pro Trp Gln Val Ser Leu Lys Gln Arg Gln Lys His Ile 75 70 Cys Gly Gly Ser Ile Val Ser Pro Gln Trp Val Ile Thr Ala Ala His 85 90 Cys Ile Ala Asn Arg Asn Ile Val Ser Thr Leu Asn Val Thr Ala Gly 100 105 Glu Tyr Asp Leu Ser Gln Thr Asp Pro Gly Glu Gln Thr Leu Thr Ile 120 125 Glu Thr Val Ile Ile His Pro His Phe Ser Thr Lys Lys Pro Met Asp 135 Tyr Asp Ile Ala Leu Leu Lys Met Ala Gly Ala Phe Gln Phe Gly His 150 155 Phe Val Gly Pro Ile Cys Leu Pro Glu Leu Arg Glu Gln Phe Glu Ala 165 170 Gly Phe Ile Cys Thr Thr 180 182

<210> 1801 <211> 202 <212> PRT

<213> Homo sapiens

<400> 1801 Met Thr Glu Ala Thr Phe Asp Thr Leu Arg Leu Trp Leu Ile Ile Leu 10 Leu Cys Ala Leu Arq Leu Ala Met Met Arg Ser His Leu Gln Ala Tyr 25 Leu Asn Leu Ala Gln Lys Cys Val Asp Gln Met Lys Lys Glu Ala Gly 40 Arg Ile Ser Thr Val Glu Leu Gln Lys Met Val Ala Arg Val Phe Tyr 55 Tyr Leu Cys Val Ile Ala Leu Gln Tyr Val Ala Pro Leu Val Met Leu 70 Leu His Thr Thr Leu Leu Leu Lys Thr Leu Gly Asn His Ser Trp Gly 90 85 Ile Tyr Pro Glu Ser Ile Ser Thr Leu Pro Val Asp Asn Ser Leu Leu 100 105 Ser Asn Ser Val Tyr Ser Glu Leu Pro Ser Ala Glu Gly Lys Met Lys 120 His Asn Ala Arg Gln Gly Pro Ala Val Pro Pro Gly Met Gln Ala Tyr 140 135 Gly Ala Ala Pro Phe Glu Asp Leu Gln Leu Asp Phe Thr Glu Met Pro 155 150 Lys Cys Gly Asp Leu Ile Pro Arg Phe Gly Leu Pro Leu Arg Ile Gly 170 Ser Asp Asn Gly Leu Ala Phe Val Ala Asp Leu Val Gln Lys Thr Ala 185 . 190 Lys Trp Lys Gly Pro Gln Ile Val Val Leu 200

<210> 1802

<211> 172 <212> PRT <213> Homo sapiens

<400> 1802 Met Asn Asn Phe Arg Ala Thr Ile Leu Phe Trp Ala Ala Ala Ala Trp 10 Ala Lys Ser Gly Lys Pro Ser Gly Glu Met Asp Glu Val Gly Val Gln 25 Lys Cys Lys Asn Ala Leu Lys Leu Pro Val Leu Glu Val Leu Pro Gly 40 Gly Gly Trp Asp Asn Leu Arg Asn Val Asp Met Gly Arg Val Met Glu 55 Leu Thr Tyr Ser Asn Cys Arg Thr Thr Glu Asp Gly Gln Tyr Ile Ile 70 75 Pro Asp Glu Ile Phe Thr Ile Pro Gln Lys Gln Ser Asn Leu Glu Met 90 85 Asn Ser Glu Ile Leu Glu Ser Trp Ala Asn Tyr Gln Ser Ser Thr Ser 105 100 Tyr Ser Ile Asn Thr Glu Leu Ser Leu Phe Ser Lys Val Asn Gly Lys 120 125 Phe Ser Thr Glu Phe Gln Arg Met Lys Thr Leu Gln Val Lys Asp Gln 135 140 Ala Ile Thr Thr Arg Val Gln Val Arg Asn Leu Val Tyr Thr Val Lys 150 155 Ile Asn Pro Thr Leu Glu Leu Ser Ser Gly Phe Arg 170

<210> 1803 <211> 158 <212> PRT <213> Homo sapiens

<400> 1803 Met Ser Leu Arg Leu Gly Pro Ala Trp Arg His Leu Thr Cys Leu Gly 10 Thr Lys His Ser Lys Ala Asn Ser Val Leu Ala Ser Gln His Ala Gly 20 25 Phe Phe Val Ala Gln Gly Arg Trp Ala Ile His Arg Ala Phe Ser Ser 40 Arg Thr Ser Pro Thr Pro Pro Arg Gly Pro Leu Leu Pro Gly Arg 55 His Pro Leu Leu Ser Arg Arg Ala Gln Ala Ile Arg Ser Ser Thr 70 Arg Pro Ser Leu Pro Ala His Leu Phe Lys Pro Ala Pro Ala Ile Ala 90 Leu Ile Val Ser Pro Leu Arg Phe Pro Arg Arg Thr Ser Pro Cys His 105 Leu Ser Gly Pro Pro Ala Pro Pro Cys Arg Thr Leu His Thr Leu Leu 120 125 Arg Pro Val Cys Val Val Arg Arg Thr Pro Pro Val Phe Phe Thr Ser 140 135 Phe Thr Pro Ala Arg Ala Ala Val Ala Ser His Pro Thr Pro 150 155

<210> 1804 <211> 102 <212> PRT <213> Homo sapiens

<400> 1804 Met Gly Leu Gly Gln Pro Gln Ala Trp Leu Leu Gly Leu Pro Thr Ala 10 Val Val Tyr Gly Ser Leu Ala Leu Phe Thr Thr Ile Leu His Asn Val 20 25 Phe Leu Leu Tyr Tyr Val Asp Thr Phe Val Ser Val Tyr Lys Ile Asn 45 40 Lys Met Ala Phe Trp Val Gly Glu Thr Val Phe Leu Leu Trp Asn Ser 60 55 Leu Asn Asp Pro Leu Phe Gly Trp Leu Ser Asp Arg Gln Phe Leu Ser 75 Ser Gln Pro Arg Ser Gly Ala Gly Leu Ser Ser Arg Ala Val Leu 85 Ala Arg Val Gln Ala Leu

<210> 1805 <211> 54 <212> PRT <213> Homo sapiens

100

102

<210> 1806 <211> 56 <212> PRT <213> Homo sapiens

<210> 1807 <211> 47 <212> PRT <213> Homo sapiens

<210> 1808 <211> 119 <212> PRT <213> Homo sapiens

<400> 1808 Met Ala Ala Ser Leu Leu Ala Val Leu Leu Leu Leu Leu Leu Glu Arg Gly Met Phe Ser Ser Pro Ser Pro Pro Pro Ala Leu Leu Glu Lys Val 2.0 25 Phe Gln Tyr Ile Asp Leu His Gln Asp Glu Phe Val Gln Thr Leu Lys 35 40 Glu Trp Val Ala Ile Glu Ser Asp Ser Val Gln Pro Val Pro Arg Phe 55 Arg Gln Glu Leu Phe Arg Met Met Ala Val Ala Ala Asp Thr Leu Gln 70 75 Arg Leu Gly Ala Arg Val Ala Ser Val Asp Met Gly Pro Gln Gln Leu 85 90 Pro Asp Gly Gln Ser Leu Pro Ile Pro Pro Val Ile Leu Ala Glu Leu 105 100 Gly Ser Asp Pro Thr Lys Gly 115 119

<210> 1809 <211> 91 <212> PRT <213> Homo sapiens

50 55 60

Arg Val Asp Val Ile Pro Leu Ser Ser Leu Gly Pro Leu Val Ser Pro
65 70 75 80

Leu Arg Cys Gln Ala Leu Pro Pro Arg Leu Ser
85 90 91

<210> 1810 <211> 58 <212> PRT <213> Homo sapiens

(213) Homo Baptem

<210> 1811 <211> 48 <212> PRT <213> Homo sapiens

<210> 1812 <211> 84 <212> PRT <213> Homo sapiens

Glu Asp Asn Phe Val Ala Leu Ala Thr Gly Gln Lys Gly Phe Gly Tyr
65 70 75 80
Lys Asn Ser *

<210> 1813

<211> 46

<212> PRT

<213> Homo sapiens

<400> 1813

Met Ala Ala Ala Asp Asp Thr Ile Leu Gly Phe Arg Ala Ala Leu Leu 1 5 10 15 15 11e Leu Val Ala Ala Ala Ala Ala Ala Leu Ser Pro Lys Val Ala Cys Arg 20 25 30 Val Gly Thr Val Arg Arg Arg Glu Thr Pro Gln Pro Ser Ala 35 40 45 46

<210> 1814

<211> 65

<212> PRT

<213> Homo sapiens

<400> 1814

 Met
 Ile
 Tyr
 Leu
 Thr
 Phe
 Pro
 Val
 Ala
 Met
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 Gln
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 Asp
 Asp
 Val
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 Lys
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 Trp
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 Arg
 Leu

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 Lys
 Leu
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 Leu
 Arg
 Asp
 Ala
 Gln
 Asn
 Ser

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<210> 1815

<211> 100

<212> PRT

<213> Homo sapiens

<400> 1815

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 Lys
 Leu
 Leu
 Asn
 Phe
 Tyr
 Ile
 Phe
 Val
 Asn
 Cys
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 Asn
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 Trp
 Ser
 Glu

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75 70 Pro Asn Ala Ile Pro Phe Ile Val Pro His Pro Gln Thr Gly Pro Asn 90 Val Arg Cys Ser 100 <210> 1816 <211> 115 <212> PRT <213> Homo sapiens <221> misc feature <222> (1)...(115) <223> Xaa = any amino acid or nothing <400> 1816 Met Phe Cys Phe Leu Val Ser Val Leu Tyr Ser Lys Ala Lys Leu Ala 10 Ser Ala Cys Gly Gly Ile Ile Tyr Phe Leu Ser Tyr Val Pro Tyr Met 20 25 Tyr Val Ala Ile Arg Glu Glu Val Ala His Asp Lys Ile Thr Ala Phe 40 Glu Lys Cys Ile Ala Ser Leu Met Ser Thr Thr Ala Phe Gly Leu Gly 55 Ser Lys Tyr Phe Ala Leu Tyr Glu Val Pro Gly Val Gly Ile Gln Trp 75 70 His Thr Phe Ser Gln Ser Pro Val Glu Gly Glu Asp Leu Asn Leu Pro 90 85 Pro Pro Pro Pro Met Met Pro Ala Pro Xaa Val Val Tyr Gly Ile Leu 105 Thr Lys * 114 <210> 1817 <211> 144 <212> PRT <213> Homo sapiens <400> 1817 Met Val Leu Gly Leu Leu Val Gln Ile Trp Ala Leu Gln Glu Ala Ser 5 Ser Leu Ser Val Gln Gln Gly Pro Asn Leu Leu Gln Val Arg Gln Gly 25

 Met
 Val
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 Ile
 Trp
 Ala
 Leu
 Gln
 Glu
 Ala
 Ser
 Gln
 Gln
 Gln
 Ile
 Trp
 Ala
 Leu
 Gln
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 Val
 Arg
 Gln
 Gln
 Gln
 Leu
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<210> 1818 <211> 115 <212> PRT <213> Homo sapiens

<400> 1818 Met Gln Ala Asp Arg Gly Gly Val Leu Phe Leu Val Ala Leu Pro Gly 10 Leu Trp Glu Thr Val Leu Arg His Pro Gly Ala Ser Pro Glu Pro Val 20 25 Ser Leu His Thr Gly Leu Ala Ala Glu Pro Leu Leu Gly Trp Arg Ala 40 Glu Val Ala Thr Ala Ala Gly Leu Gln Asp Arg Arg Ile Gly Arg Arg 55 60 Ser Leu Pro Ala Thr Leu Pro Pro Pro Phe Pro Gln Ala Gly Asp Leu 70 75 Arg Glu Ser Ile Leu Leu Pro Cys Arg Glu Ser Arg Ser Thr Ser 90 Trp Leu Ser Pro Tyr Trp Val Pro Glu Ile Pro Gly Thr Leu His Asp 105 Arg Gly Arg 115

<210> 1819 <211> 70 <212> PRT <213> Homo sapiens

<210> 1820 <211> 635 <212> PRT <213> Homo sapiens

<pre><400> 1820 Met Leu Arg Ser Leu Leu Val Tyr Met Leu Phe Leu Leu Val Thr Le</pre>															
1				5					10					15	
			20					25					30	Arg	
		35					40					45		Ala	
	50					55					60			Leu	
65					70					75				Arg	80
				85					90					Gly 95	
			100					105					110	Asp	
_		115					120					125		Ala	
	130					135					140			Ala	
145					150					155				Glu	160
	_	_		165					170					175	
_		_	180				•	185					190	Ala	
		195					200					205		Ala	
	210					215					220			Arg	
225					230					235				Leu	240
				245					250					Arg 255	
			260					265					270	Leu	
		275					280					285		Leu	
	290					295					300			Arg	
305					310					315				Arg	320
				325					330					Gln 335	
			340			•		345					350		Arg
		355					360					365		Leu	
	370	_				375					380			Val	
385		_			390					395		•			Thr 400
				405					410					Pro 415	
			420					425					430		Leu
Gly	Ala	Val 435		Leu	Arg	Trp	Arg 440		His	Ala	Leu	Arg 445		Glu	Leu

Tyr Arg Pro Ala Trp Glu Pro Gln Asp Tyr Glu Met Val Glu Leu Phe 455 Leu Arg Arg Leu Arg Leu Trp Met Gly Leu Ser Lys Val Lys Glu Phe 475 470 Arg His Lys Val Arg Phe Glu Gly Met Glu Pro Leu Pro Ser Arg Ser 490 485 Ser Arg Gly Ser Lys Val Ser Pro Asp Val Pro Pro Pro Ser Ala Gly 505 Ser Asp Ala Ser His Pro Ser Thr Ser Ser Gln Leu Asp Gly Leu 520 Ser Val Ser Leu Gly Arg Leu Gly Thr Arg Cys Glu Pro Glu Pro Ser Arg Leu Gln Ala Val Phe Glu Ala Leu Leu Thr Gln Phe Asp Arg Leu 550 555 Asn Gln Ala Thr Glu Asp Val Tyr Gln Leu Glu Gln Gln Leu His Ser 570 565 Leu Gln Gly Arg Arg Ser Ser Arg Ala Pro Ala Gly Ser Ser Arg Gly 585 580 Pro Ser Pro Gly Leu Arg Pro Ala Leu Pro Ser Arg Leu Ala Arg Ala 605 600 Ser Arg Gly Val Asp Leu Ala Thr Gly Pro Ser Arg Thr Pro Leu Arg 615 Ala Lys Asn Lys Val His Pro Ser Ser Thr * 630

<210> 1821 <211> 84 <212> PRT

<213> Homo sapiens

<210> 1822 <211> 108 <212> PRT <213> Homo sapiens

84

<210> 1823 <211> 74 <212> PRT <213> Homo sapiens

<210> 1824 <211> 58 <212> PRT <213> Homo sapiens

<210> 1825 <211> 225 <212> PRT <213> Homo sapiens

<400> 1825

Met Ala Cys Lys Gly Leu Leu Gln Gln Val Gln Gly Pro Arg Leu Pro Trp Thr Arg Leu Leu Leu Leu Leu Val Phe Ala Val Gly Phe Leu 25 Cys His Asp Leu Arg Ser His Ser Ser Phe Gln Ala Ser Leu Thr Gly Arg Leu Leu Arg Ser Ser Gly Phe Leu Pro Ala Ser Gln Gln Ala Cys 55 Ala Lys Leu Tyr Ser Tyr Ser Leu Gln Gly Tyr Ser Trp Leu Gly Glu 70 Thr Leu Pro Leu Trp Gly Ser His Leu Leu Thr Val Val Arg Pro Ser 90 8.5 Leu Gln Leu Ala Trp Ala His Thr Asn Ala Thr Val Ser Phe Leu Ser 105 100 Ala His Cys Ala Ser His Leu Ala Trp Phe Gly Asp Ser Leu Thr Ser 125 120 Leu Ser Gln Arg Leu Gln Ile Gln Leu Pro Asp Ser Val Asn Gln Leu 140 135 Leu Arg Tyr Leu Arg Glu Leu Pro Leu Leu Phe His Gln Asn Val Leu 150 155 Leu Pro Leu Trp His Leu Leu Leu Glu Ala Leu Ala Trp Ala Gln Glu 170 His Cys His Glu Ala Cys Arg Gly Glu Val Thr Trp Asp Cys Met Lys 185 Thr Gln Leu Ser Glu Ala Val His Trp Thr Trp Leu Cys Leu Gln Asp 205 200 Ile Thr Val Ala Phe Leu Asp Trp Ala Leu Ala Leu Ile Ser Gln Gln 215

<210> 1826 <211> 119 <212> PRT <213> Homo sapiens

<400> 1826 Met Tyr Arg Glu Val Cys Ser Ile Arg Phe Leu Phe Thr Ala Val Ser 10 Leu Leu Ser Leu Phe Leu Ser Ala Phe Trp Leu Gly Leu Leu Tyr Leu 25 Val Ser Pro Leu Glu Asn Glu Pro Lys Glu Met Leu Thr Leu Ser Glu Tyr His Glu Arg Ala Arg Ser Gln Gly Gln Gln Leu Leu Gln Phe Gln Ala Glu Leu Asp Lys Leu His Lys Glu Ala Ser Leu Val Cys Gly Cys 75 Pro Ser Leu Arg Glu Val Pro Ser Ser Ala Val Ser Arg Leu Glu Pro 90 Pro Ser Ile Ala Gln Pro Leu Leu Ser Arg Leu Gln Leu Tyr Leu Ser 100 110 105 Asp Pro Ser Ser Tyr Leu Val 115

<210> 1827 <211> 58 <212> PRT <213> Homo sapiens

<400> 1827

 Met
 Lys
 Leu
 Met
 Arg
 Pro
 Leu
 Met
 Leu
 Tyr
 Ile
 Ser
 Gln
 Leu
 Tyr

 Met
 Leu
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 Lys
 Arg
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 Leu
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<210> 1828 <211> 102 <212> PRT <213> Homo sapiens

<400> 1828

Met Gln Pro Ser Gly Leu Glu Gly Pro Gly Thr Phe Gly Arg Trp Pro 5 10 Leu Leu Ser Leu Leu Leu Leu Leu Leu Leu Gln Pro Val Thr Cys 25 Ala Tyr Thr Thr Pro Gly Pro Pro Arg Ala Leu Thr Thr Leu Gly Ala 40 Pro Arg Ala His Thr Met Pro Gly Thr Tyr Ala Pro Ser Thr Thr Leu Ser Ser Pro Ser Thr Gln Gly Leu Gln Glu Gln Ala Arg Ala Leu Met 70 75 Arg Asp Phe Pro Leu Val Asp Gly His Asn Asp Leu Pro Leu Val Leu 90 85 Arg Gln Val Tyr His Asn 100

<210> 1829 <211> 88 <212> PRT <213> Homo sapiens

<400> 1829

 Met
 Arg
 Lys
 Ile
 Tyr
 Thr
 Thr
 Val
 Leu
 Phe
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 Asn
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 Glu
 Gln
 Trp

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 Leu
 Ile

 Val
 Ala
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 Ile
 Leu
 Ser
 Trp
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 Pro
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 Trp
 Thr
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Met Leu Ser Asp Tyr Ala Lys Pro 85 88

<210> 1830

<211> 120

<212> PRT

<213> Homo sapiens

<400> 1830

Met Lys Trp Arg Arg Lys Ser Ala Tyr Trp Lys Ala Leu Lys Val Phe 10 Lys Leu Pro Val Glu Phe Leu Leu Leu Thr Val Pro Val Val Asp 20 25 Pro Asp Lys Asp Asp Gln Asn Trp Lys Arg Pro Leu Asn Cys Leu His 40 Leu Val Ile Ser Pro Leu Val Val Leu Thr Leu Gln Ser Gly Thr 55 Tyr Gly Val Tyr Glu Ile Gly Gly Leu Val Pro Val Trp Val Val Val 70 75 Val Ile Ala Gly Thr Ala Leu Ala Ser Val Thr Phe Phe Ala Thr Ser 85 90 Asp Ser Gln Pro Pro Arg Leu His Trp Leu Phe Ala Phe Leu Gly Phe 105

<210> 1831

115

<211> 64

<212> PRT

<213> Homo sapiens

Leu Thr Ser Ala Leu Trp Ile Asn

<400> 1831

<210> 1832

<211> 89

<212> PRT

<213> Homo sapiens

<400> 1832

Met Gly Ile Lys His Phe Ser Gly Leu Phe Val Leu Leu Cys Ile Gly 1 5 10 15 Phe Gly Leu Ser Ile Leu Thr Thr Ile Gly Glu His Ile Val Tyr Arg

<210> 1833 <211> 60 <212> PRT <213> Homo sapiens

<210> 1834 <211> 62 <212> PRT <213> Homo sapiens

<210> 1835 <211> 71 <212> PRT <213> Homo sapiens

<210> 1836 <211> 110 <212> PRT <213> Homo sapiens

<400> 1836 Met Leu Met Tyr Met Phe Tyr Val Leu Pro Phe Cys Gly Leu Ala Ala 5 10 Tyr Ala Leu Thr Phe Pro Gly Cys Ser Trp Leu Pro Asp Trp Ala Leu 20 25 Val Phe Ala Gly Gly Ile Gly Gln Ala Gln Phe Ser His Met Gly Ala 40 Ser Met His Leu Arg Thr Pro Phe Thr Tyr Arg Val Pro Glu Asp Thr 55 Trp Gly Cys Phe Phe Val Cys Asn Leu Leu Tyr Ala Leu Gly Pro His 70 75 Leu Leu Ala Tyr Arg Cys Leu Gln Trp Pro Ala Phe Phe His Gln Pro 90 Pro Pro Ser Asp Pro Leu Ala Leu His Lys Lys Gln His

105

<210> 1837 <211> 91 <212> PRT <213> Homo sapiens

<210> 1838 <211> 201 <212> PRT <213> Homo sapiens

<400> 1838 Met Pro Ile Gly Leu Arg Gly Leu Met Ile Ala Val Met Leu Ala Ala 10 Leu Met Ser Ser Leu Thr Ser Ile Phe Asn Ser Ser Ser Thr Leu Phe 20 25 Thr Met Asp Ile Trp Arg Arg Leu Arg Pro Arg Ser Gly Glu Arg Glu 40 Leu Leu Leu Val Gly Arg Leu Val Ile Val Ala Leu Ile Gly Val Ser Val Ala Trp Ile Pro Val Leu Gln Asp Ser Asn Ser Gly Gln Leu Phe 70 75 Ile Tyr Met Gln Ser Val Thr Ser Ser Leu Ala Pro Pro Val Thr Ala 90 85 Val Phe Val Leu Gly Val Phe Trp Arg Arg Ala Asn Glu Gln Gly Ala 100 105 Phe Trp Gly Leu Ile Ala Gly Leu Val Val Gly Ala Thr Arg Leu Val 125 120 Leu Glu Phe Leu Asn Pro Ala Pro Pro Cys Gly Glu Pro Asp Thr Arg 135 140 Pro Ala Val Leu Gly Ser Ile His Tyr Leu His Phe Ala Val Ala Leu 155 150 Phe Ala Leu Ser Gly Ala Val Val Ala Gly Ser Leu Leu Thr Pro 170 Pro Pro Gln Ser Val Gln Ile Glu Asn Leu Thr Trp Trp Thr Leu Ala 185 Gln Asp Val Pro Leu Gly Thr Lys Ala 200 201

<210> 1839 <211> 130

<212> PRT

<213> Homo sapiens

<221> misc_feature

<222> (1)...(130)

<223> Xaa = any amino acid or nothing

<400> 1839 Met Leu Phe Phe Leu Gln Ser Leu Phe Met Leu Ala Thr Val Val Leu 10 1 5 Tyr Phe Ser His Leu Lys Glu Tyr Val Ala Ser Met Val Phe Ser Leu 25 Ala Leu Gly Trp Thr Asn Met Leu Tyr Tyr Thr Arg Gly Phe Gln Gln 40 45 Met Gly Ile Tyr Ala Val Met Ile Glu Lys Met Ile Leu Arg Asp Leu Cys Arg Phe Met Phe Val Tyr Ile Val Phe Leu Phe Gly Phe Ser Thr 75 · Ala Val Val Thr Leu Ile Glu Asp Gly Lys Asn Asp Ser Leu Pro Ser 85 Glu Ser Thr Ser His Arg Trp Arg Gly Phe Ser Xaa Thr Pro Leu Xaa 105 Leu Leu His Lys Leu Tyr Ser Thr Cys Leu Glu Leu Ser Asn Ser Thr 120

Xaa Asp 130

<210> 1840

<211> 47

<212> PRT

<213> Homo sapiens

<400> 1840

Met Asn Arg Val Met Arg Gly Leu Ala Ile Thr Thr Cys Leu Leu Ser Met Leu Gln Ala Ile Thr Ile Ser Pro Ser Ile Leu Trp Asn His 20 25 Ala Ala Val Gln Tyr Val His Gly His Ser Leu Val Gln Ala 40

<210> 1841

<211> 82

<212> PRT

<213> Homo sapiens

<400> 1841

Met Thr Ala Arg Leu Met Arg Ser Leu Leu Ala Ala Gln Leu Thr Phe 10 Val Tyr Arg Val Ala His Leu Met Asn Val Ala Gln Arg Ile Arg Gly 25 Asn Arg Pro Ile Lys Asn Glu Arg Leu Leu Ala Leu Leu Gly Asp Asn 40 Glu Lys Met Asn Leu Ser Asp Val Glu Leu Ile Pro Leu Pro Leu Glu 55 Pro Gln Val Lys Ile Arg Gly Ile Ile Pro Glu Thr Ala Thr Leu Phe Lys Ser

82

<210> 1842

<211> 77

<212> PRT

<213> Homo sapiens

<400> 1842

Met Val Ala Asn Met Phe Tyr Ile Val Val Ile Met Ala Leu Val Leu 10 Leu Ser Phe Gly Val Pro Arg Lys Ala Ile Leu Tyr Pro His Glu Ala 20 25 Pro Ser Trp Thr Leu Ala Lys Asp Ile Val Phe His Pro Tyr Trp Met 40 Ile Phe Gly Glu Val Tyr Ala Tyr Glu Ile Asp Val Cys Ala Asn Asp Ser Val Ile Pro Gln Ile Cys Gly Pro Ser Thr Arg Pro

65 70 75 77

<210> 1843 <211> 109 <212> PRT <213> Homo sapiens

<400> 1843 Met Met His Asn Ile Ile Val Lys Glu Leu Ile Val Thr Phe Phe Leu 10 5 Gly Ile Thr Val Val Gln Met Leu Ile Ser Val Thr Gly Leu Lys Gly 20 25 Val Glu Ala Gln Asn Gly Ser Glu Ser Glu Val Phe Val Gly Lys Tyr 40 45 Glu Thr Leu Val Phe Tyr Trp Pro Ser Leu Leu Cys Leu Ala Phe Leu 55 60 Leu Gly Arg Phe Leu His Met Phe Val Lys Ala Leu Arg Val His Leu 70 75 Gly Trp Glu Leu Gln Val Glu Glu Lys Ser Val Leu Glu Val His Gln 90 Gly Glu His Val Lys Gln Leu Leu Arg Ile Pro Arg Pro

105

<210> 1844 <211> 85 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(85) <223> Xaa = any amino acid or nothing

<210> 1845 <211> 110 <212> PRT <213> Homo sapiens

<400> 1845 Met Tyr Ala Leu Tyr Ile Thr Val His Gly Tyr Phe Leu Ile Thr Phe 10 Leu Phe Gly Met Val Val Leu Ala Leu Val Val Trp Lys Ile Phe Thr 20 Leu Ser Arg Ala Thr Ala Val Lys Glu Arg Gly Lys Asn Arg Lys Lys 40 Val Leu Thr Leu Leu Gly Leu Ser Ser Leu Val Gly Val Thr Trp Gly 55 Leu Ala Ile Phe Thr Pro Leu Gly Leu Ser Thr Val Tyr Ile Phe Ala 70 75 Leu Phe Asn Ser Leu Gln Gly Val Phe Ile Cys Cys Trp Phe Thr Ile 85 90 Leu Tyr Leu Pro Ser Gln Ser Thr Thr Val Ser Ser Ser Thr 105 100

<210> 1846 <211> 94 <212> PRT <213> Homo sapiens

<210> 1847 <211> 1300 <212> PRT <213> Homo sapiens

				85					90					9.5	
Cys	Pro	Asp	Tyr		Ser	Phe	Cys	Ala 105		Val	His	Asn		95 Thr	Ser
Pro	Pro	Ser 115	Ser	Lys	Lys	Ala	Pro 120		Pro	Ser	Gly	Ala 125	110 Ser	Gln	Thr
Ile	Lys 130	-	Thr	Thr	Lys	Arg 135	Ser	Pro	Lys	Pro	Pro 140		Lys	Lys	Lys
Thr 145	Lys	Lys	Val	Ile	Glu 150			Glu	Ile	Thr 155		Glu	His	Ser	Val 160
Ser	Glu	Asn	Gln	Glu 165	Ser	Ser	Ser	Ser	Ser 170	Ser	Ser	Ser	Ser	Ser 175	Ser
			Trp 180					185					190		_
		195	Lys				200					205			
	210		Pro			215					220			-	
225			Asn		230					235					240
			Asn Pro	245					250					255	_
			260 Leu					265					270		
		275	Thr				280					285		_	
	290		Lys			295					300				
305			Thr		310					315					320
Thr	Thr	Thr	Lys	325 Gly	Pro	Ala	Leu		330 Thr	Pro	Lys	Glu		335 Thr	Pro
Thr	Thr	Pro	340 Lys	Glu	Pro	Ala		345 Thr	Thr	Pro	Lys		350 Pro	Thr	Pro
Thr	Thr 370		Lys	Ser	Ala	Pro 375	360 Thr	Thr	Pro	Lys	Glu 380	365 Pro	Ala	Pro	Thr
Thr 385		Lys	Ser	Ala	Pro 390		Thr	Pro	Lys	Glu 395		Ala	Pro	Thr	Thr 400
Thr	Lys	Glu	Pro	Ala 405		Thr	Thr	Pro	Lys 410		Pro	Ala	Pro	Thr 415	
			Pro 420					425					430		
		435	Ala				440	=	_			445			
	450		Ala			455					460				
465			Ala		470		_			475					480
			Pro Pro	485					490					495	
			500 Pro					505		,		•	510		
		515	Thr				520					525			
	530		Thr			535					540				
545					550					555			-		560

Ala	Pro	Thr	Thr	Pro 565	Lys	Glu	Pro	Ala	Pro 570	Thr	Thr	Pro	Lys	Lys 575	Pro
Ala	Pro	Thr	Thr 580	Pro	Lys	Glu	Pro	Ala 585	Pro	Thr	Thr	Pro	Lys 590	Glu	Pro
Ala	Pro	Thr 595	Thr	Thr	Lys	Lys	Pro 600		Pro	Thr	Ala	Pro 605	Lys	Glu	Pro
Ala	Pro 610		Thr	Pro	Lys	Glu 615		Ala	Pro	Thr	Thr 620		Lys	Lys	Leu
		Thr	Thr	Pro			Leu	Ala	Pro	Thr 635		Pro	Glu	Lys	Pro 640
625 Ala	Pro	Thr	Thr		630 Glu	Glu	Leu	Ala			Thr	Pro	Glu		
Thr	Pro	Thr	Thr	645 Pro	Glu	Glu	Pro		650 Pro	Thr	Thr	Pro		655 Ala	Ala
Ala	Pro	Asn	660 Thr	Pro	Lys	Glu	Pro	665 Ala	Pro	Thr	Thr	Pro	670 Lys	Glu	Pro
		675					680					685			
Ala	Pro 690	Thr	Thr	Pro	Lys	Glu 695	Pro	Ala	Pro	Thr	700	Pro	ьуs	GIU	Thr
705			Thr		710					715					720
Ala	Pro	Thr	Thr	Pro 725	Lys	Lys	Pro	Ala	Pro 730	Lys	Glu	Leu	Ala	Pro 735	Thr
Thr	Thr	Lys	Glu 740	Pro	Thr	Ser	Thr	Thr 745	Ser	Asp	Lys	Pro	Ala 750	Pro	Thr
Thr	Pro	Lys 755	Gly	Thr	Ala	Pro	Thr 760	Thr	Pro	Lys	Glu	Pro 765	Ala	Pro	Thr
Thr	Pro 770	Lys	Glu	Pro	Ala	Pro 775	Thr	Thr	Pro	Lys	Gly 780	Thr	Ala	Pro	Thr
Thr 785	Leu	Lys	Glu	Pro	Ala 790	Pro	Thr	Thr	Pro	Lys 795	Lys	Pro	Ala	Pro	Lys 800
	Leu	Ala	Pro	Thr 805		Thr	Lys	Gly	Pro 810	Thr	Ser	Thr	Thr	Ser 815	Asp
Lys	Pro	Ala	Pro 820		Thr	Pro	Lys	Glu 825	Thr	Ala	Pro	Thr	Thr 830	Pro	Lys
Glu	Pro	Ala 835	Pro	Thr	Thr	Pro	Lys 840		Pro	Ala	Pro	Thr 845	Thr	Pro	Glu
Thr	Pro 850		Pro	Thr	Thr	Ser 855			Ser	Thr	Pro 860	Thr	Thr	Thr	Lys
	Pro	Thr	Thr	Ile			Ser	Pro	Asp	Glu 875	Ser	Thr	Pro	Glu	Leu 880
865 Ser		Glu	Pro	Thr	870 Pro	Lys	Ala			Asn	Ser				Pro
Gly	Val	Pro	Thr	885 Thr	Lys	Thr	Pro	Ala	890 Ala		Lys		Glu	895 Met	Thr
Thr	Thr	Ala	900 Lvs	Asp	Lvs	Thr	Thr	905 Glu	Arq	Asp	Leu	Arg	910 Thr	Thr	Pro
		915					920					925			Thr
	930					935					940				
Thr 945	Glu	Lys	Thr	Thr	Glu 950	Ser	Lys	Ile	Thr	955	Thr	Thr	Tnr	GIR	Val 960
	Ser	Thr	Thr	Thr 965	Gln	Asp	Thr	Thr	Pro 970		Lys	Ile	Thr	Thr 975	Leu
Lys	Thr	Thr	Thr 980		Ala	Pro	Lys	Val 985			Thr	Lys	Lys 990		Ile
Thr	Thr	Thr	Glu	Ile	Met		Lys 1000	Pro	Glu	Glu		Ala 1005	Lys		Lys
_		Ala		Asn		Lys	Ala		Thr		Lys			Lys	Pro
	1010 Lys		Pro	Lys		1015 Pro		Ser	Thr		1020 Lys	Pro	Lys	Thr	Met

1030 1035 Pro Arg Val Arg Lys Pro Lys Thr Thr Pro Thr Pro Arg Lys Met Thr 1045 1050 Ser Thr Met Pro Glu Leu Asn Pro Thr Ser Arg Ile Ala Glu Ala Met 1060 1065 1070 Leu Gln Thr Thr Arg Pro Asn Gln Thr Pro Asn Ser Lys Leu Val 1075 1080 1085 Glu Val Asn Pro Lys Ser Glu Asp Ala Gly Gly Ala Glu Gly Glu Thr 1090 1095 1100 Pro His Met Leu Leu Arg Pro His Val Phe Met Pro Glu Val Thr Pro 1105 1110 1115 1120 Asp Met Asp Tyr Leu Pro Arg Val Pro Asn Gln Gly Ile Ile Ile Asn 1125 1130 1135 Pro Met Leu Ser Asp Glu Thr Asn Ile Cys Asn Gly Lys Pro Val Asp 1140 1145 1150 Gly Leu Thr Thr Leu Arg Asn Gly Thr Leu Val Ala Phe Arg Gly His 1155 1160 1165 Tyr Phe Trp Met Leu Ser Pro Phe Ser Pro Pro Ser Pro Ala Arg Arg 1170 1175 1180 Ile Thr Glu Val Trp Gly Ile Pro Ser Pro Ile Asp Thr Val Phe Thr 1185 1190 1195 Arg Cys Asn Cys Glu Gly Lys Thr Phe Phe Lys Asp Ser Gln Tyr 1205 1210 1215 Trp Arg Phe Thr Asn Asp Ile Lys Asp Ala Gly Tyr Pro Lys Pro Ile 1220 1225 Phe Lys Gly Phe Gly Gly Leu Thr Gly Gln Ile Val Ala Ala Leu Ser 1235 1240 1245 Thr Ala Lys Tyr Lys Asn Trp Pro Glu Ser Val Tyr Phe Phe Lys Arg 1255 1260 Gly Ser Ile Gln Gln Tyr Ile Tyr Lys Gln Glu Pro Val Gln Lys 1265 1270 1275 Cys Pro Gly Arg Arg Pro Ala Leu Asn Tyr Pro Val Tyr Gly Glu Thr 1285 1290 Asp Thr Gly * . 1299

<210> 1848 <211> 103

<212> PRT

<213> Homo sapiens

<400> 1848

Met Asn Pro Ala Val Arg Gln Arg Cys Leu Leu Phe Cys Phe Gln Gln 1 5 10 Lys Leu Ile Leu Ser His Phe Phe Leu Leu Gln Val Pro Gln Trp Cys Ala Glu Tyr Cys Leu Ser Ile His Tyr Gln His Gly Gly Val Ile Cys 35 40 Thr Gln Val His Lys Gln Thr Val Val Gln Leu Ala Leu Arg Val Ala 55 60 Asp Glu Met Asp Val Asn Ile Gly His Glu Val Gly Tyr Val Ile Pro 70 75 Phe Glu Asn Cys Cys Thr Asn Glu Thr Ile Leu Arg Leu Val Cys Gly 85 90 Val Gln Ser Ala Pro Cys * 100 102

<210> 1849 <211> 50 <212> PRT <213> Homo sapiens

<210> 1850 <211> 84 <212> PRT <213> Homo sapiens

<210> 1851 <211> 51 <212> PRT <213> Homo sapiens

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<210> 1852
<211> 54
<212> PRT
<213> Homo sapiens
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<210> 1853 <211> 129 <212> PRT <213> Homo sapiens

<400> 1853 Met Ala Val Val Arg Val Met Val Val Arg Val Thr Ala Val Val 1 5 10 Arg Val Met Val Val Val Val Val Val Arg Val Met Val Val 20 25 Val Arg Ile Thr Ala Val Leu Arg Val Met Val Val Arg Ile Met 40 Ala Val Ile Arg Val Met Val Val Val Arg Val Thr Ala Ile Val Gly . 55 Val Met Val Val Ile Arg Val Thr Ala Ile Val Ser Ile Met Val Val 70 Val Arg Val Met Val Val Val Arg Val Met Val Val Ala Arg Pro Met 90 Val Val Val Arg Val Met Ala Val Val Arg Val Met Ala Asp Ser Ala 100 105 Leu Arg Ala Ile Cys Ser Ser Ser Leu Asn Val Thr Phe Ser Leu Glu

<210> 1854
<211> 190
<212> PRT
<213> Homo sapiens
<221> misc_feature
<222> (1)...(190)
<223> Xaa = any amino acid or nothing

<400> 1854

PCT/US01/02687 WO 01/54477

Met Ser Cys Phe Gly Leu Leu Leu Gly Gly Leu Thr Pro Arg Val Leu Ser Thr Glu Glu Gln Leu Pro Pro Gly Phe Pro Ser Ile Asp Met Gly 25 Pro Gln Leu Lys Val Val Glu Lys Ala Arg Thr Ala Thr Met Leu Cys Ala Ala Gly Gly Asn Pro Asp Pro Glu Ile Ser Trp Phe Lys Asp Phe 55 Leu Pro Val Asp Pro Ala Thr Ser Asn Gly Arg Ile Lys Gln Leu Arg 70 Ser Gly Glu Gln Arg Ala Gly Val Lys Gly Pro Cys Arg Pro Gln Asn 8.5 90 Lys Arg Leu Val Arg Ser Gln His Ser Leu Leu Pro Trp Ala Trp Ala 100 105 Pro Pro Gly Leu Ser Gly Gly Tyr Leu Val Gly Trp Ala Gly Ser Tyr 120 125 Cys Arg Cys Ala Trp Leu Arg Glu Glu Ser Ser Trp Leu Ala Val Pro 135 140 Leu Pro Ser Ser Asp Cys Gln Thr Pro Asp Phe Gly Pro Val Leu Pro 150 155 Leu Pro Ala His Val Met Cys Gln Cys Gly Gly Leu Phe Lys Gly Ala 170 Leu Trp Met Leu Thr Leu Leu Pro Cys Xaa Leu Ala * 185

<210> 1855 <211> 78 <212> PRT

<213> Homo sapiens

<400> 1855 Met Val Val Ser Ala Trp Ile Gly Leu Glu Ala Thr Val Val Ala Ala 10 5 Cys Leu Ala Leu Leu Gly Ser Val Val Arg Glu Thr Ser Thr Ser Ala 25 Ser Pro Thr Pro Ala Ala Leu Arg Ala Ala Trp Thr Val Tyr Ser Ser 40 Pro Met Thr Thr Cys Val Phe Ala Val Val Pro Leu Leu Ala Gly Thr 55 60 Val Lys Pro Ser Ser Met Cys Val Pro Arg Cys Pro Ala 70

<210> 1856 <211> 67 <212> PRT <213> Homo sapiens

<400> 1856 Met Thr Asn Trp Met Leu Leu Leu Ala Ser Arg Ile Phe Gln Ser Leu 10 Ala Ile Pro Lys Gln Leu Gly Leu Arg Arg Glu Met Pro Ser Gly Ser 20 25 Pro Thr Thr Asn Ser Ser Ser Gly Cys Ile Arg Asn Leu Glu Tyr Ser

35 40 45
Thr Leu Met Gly Ser Glu Met Pro Met Ala Leu Ala Ala Glu Thr Trp
50 55 60
Leu Leu *
65 66

<210> 1857 <211> 107 <212> PRT <213> Homo sapiens

1200 110000 00021011

<400> 1857 Met Leu Leu Met Phe Leu Leu Ala Thr Cys Leu Leu Ala Ile Ile Phe 1 5 10 Val Pro Gln Glu Met Gln Thr Leu Arg Val Val Leu Ala Thr Leu Gly 20 25 Val Gly Ala Ala Ser Leu Gly Ile Thr Cys Ser Thr Ala Gln Glu Asn 40 Glu Leu Ile Pro Ser Ile Ile Arg Gly Arg Ala Thr Gly Ile Thr Gly 55 Asn Phe Ala Asn Ile Gly Gly Ala Leu Ala Ser Leu Val Met Ile Leu 70 75 Ser Ile Tyr Ser Arg Pro Leu Pro Trp Ile Ile Tyr Gly Val Phe Ala 85 Ile Leu Ser Gly Leu Val Val Leu Leu Pro

<210> 1858 <211> 134 <212> PRT <213> Homo sapiens

<400> 1858 Met Ile Pro Pro Ala Ile Phe Trp Val Leu Ile Ile Phe Gly Trp Thr 10 Leu Val Tyr Gly Phe Val Tyr Phe Thr Thr Gly Glu Thr Ile Met Asp 20 25 Lys Leu Leu Arg Val Leu Tyr Trp Ile Leu Val Lys Thr Phe Phe Arg 40 Glu Ile Ser Val Ser His Gln Glu Arg Ile Pro Lys Asp Lys Pro Val 55 Met Leu Val Cys Ala Pro His Ala Asn Gln Phe Val Asp Gly Met Val 75 Ile Ser Thr His Leu Asp Arg Lys Val Tyr Phe Val Gly Ala Ala Ser 90 Ser Phe Arg Lys Tyr Lys Val Val Gly Leu Phe Met Lys Leu Met Ala 110 105 Ser Ile Ile Ser Gly Glu Arg His Gln Asp Val Lys Lys Val Leu Thr 120 Gly Met Ala Thr Glu Lys 134

<210> 1859 <211> 82 <212> PRT <213> Homo sapiens

<210> 1860 <211> 46 <212> PRT <213> Homo sapiens

<210> 1861 <211> 128 <212> PRT <213> Homo sapiens

100 105 110

Gly Ile Tyr Phe Leu Gly Gln Ala His Val Ile Ser Lys Leu Asn Met
115 120 125 128

<210> 1862 <211> 58 <212> PRT

<213> Homo sapiens

<400> 1862

 Met
 Trp
 Asp
 Met
 Leu
 Pro
 Trp
 Gly
 Ile
 Thr
 Trp
 Val
 Leu
 Leu
 Leu
 Trp
 Val
 Ile
 Gly
 Phe
 Thr
 Trp
 Val

 Cys
 Lys
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 Asp
 Arg
 Asp
 Ser
 Try
 Leu
 Glu
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 Asp
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<210> 1863 <211> 50 <212> PRT <213> Homo sapiens

<210> 1864 <211> 90 <212> PRT <213> Homo sapiens

Gly Val Glu Leu Leu Val Cys Ser Pro Leu Glu Ala Leu Gly Pro Leu 65 70 75 80

Leu Cys Leu Gly Glu Leu Gly Leu Gln Ala 85 90

<210> 1865 <211> 125 <212> PRT <213> Homo sapiens

<400> 1865 Met Arg Leu Gly Leu Leu Leu Ala Arg His Trp Cys Ile Ala Gly Val Phe Pro Gln Lys Phe Asp Gly Asp Ser Ala Tyr Val Gly Met Ser 20 25 Asp Gly Asn Pro Glu Leu Leu Ser Thr Ser Gln Thr Tyr Asn Gly Gln 40 Ser Glu Asn Asn Glu Asp Tyr Glu Ile Pro Pro Ile Thr Pro Pro Asn 55 60 Leu Pro Glu Pro Ser Leu Leu His Leu Gly Asp His Glu Ala Ser Tyr 75 70 His Ser Leu Cys His Gly Leu Thr Pro Asn Gly Leu Leu Pro Ala Tyr 90 85 Ser Tyr Gln Ala Met Asp Leu Pro Ala Ile Met Val Ser Asn Met Leu 105 100 Ala Gln Asp Ser His Leu Leu Ser Gly Gln Leu Pro Thr 115 120

<210> 1866 <211> 129 <212> PRT <213> Homo sapiens

<400> 1866 Met Cys Phe Leu Asn Lys Leu Leu Leu Leu Ala Ala Leu Asp Trp Leu 10 Phe Gln Ile Pro Thr Val Pro Glu Asp Leu Phe Phe Leu Glu Glu Gly 20 25 Pro Ser Tyr Ala Phe Glu Val Asp Thr Val Ala Pro Glu His Gly Leu 40 Asp Asn Ala Pro Val Val Asp Gln Gln Leu Leu Tyr Thr Cys Cys Pro 55 Tyr Ile Gly Glu Leu Arg Lys Leu Leu Ala Ser Trp Val Ser Gly Ser 75 Ser Gly Arg Ser Gly Gly Phe Met Arg Lys Ile Thr Pro Thr Thr 90 Thr Ser Leu Gly Ala Gln Pro Ser Gln Thr Ser Gln Gly Leu Gln Ala 105 Gln Leu Ala Gln Ala Phe Phe His Asn Gln Pro Pro Ser Leu Arg Arg 125 120 Thr 129

<210> 1867 <211> 80 <212> PRT <213> Homo sapiens

<400> 1867

 Met
 Met
 Arg
 Leu
 Glu
 Lys
 Phe
 Val
 Trp
 Ser
 Val
 Met
 Ala
 Leu
 Gly

 Trp
 Phe
 Val
 Phe
 Arg
 Gln
 Gln
 Asn
 Cys
 Trp
 Ala
 Leu
 Trp
 Ser
 Lys
 Ser

 Val
 Leu
 Ile
 Ser
 Trp
 Ser
 Arg
 Pro
 Leu
 Thr
 Arg
 Ser
 Met
 Ser
 Asp
 Leu

 Arg
 Arg
 Lys
 Arg
 Pro
 Leu
 Thr
 Arg
 Ser
 Glu
 Leu
 Tyr
 Ser
 Ser
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 Lys
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 Glu
 Arg
 Ala
 Leu
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<210> 1868 <211> 113 <212> PRT <213> Homo sapiens

(213) HOMO Sapiens

 Ala Ser
 Arg
 Thr
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 Arg
 Trp
 Pro
 Ala
 Leu
 Gly
 Ala

 Bet
 Leu
 Val
 Tyr
 Gly
 Thr
 Ile
 Arg
 Trp
 Pro
 Ala
 Leu
 Gly
 Ala

 Pro
 Arg
 Trp
 Trp
 Pro
 Pro
 Pro
 Gly
 Val
 Trp
 Ser
 Gly
 Ile

 Arg
 Thr
 Pro
 Arg
 Ala
 Arg
 Ser
 Leu
 Arg
 Gly
 Thr
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 Arg
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<210> 1869 <211> 72 <212> PRT <213> Homo sapiens

<400> 1869
Met Phe Leu Trp Val Lys Arg Leu Leu Phe Ala Ala Ser Leu Leu Ala
1 5 10 15

 Ser Asp Ser Ser Thr Ile Leu Cys Ser Arg Asp Leu Ile Leu Glu Ser 25
 30

 Ile Ala Leu Ile Ile Ala Phe Cys Ser Leu Arg Ile Leu Pro Phe Ser 35
 40

 Trp Ala Ser Ser Ser Cys Leu Cys Ile Met Phe Ser Ser Val Ser Leu 50
 55

 Ser Ala Arg Ser Phe Phe Ile *
 65

<210> 1870 <211> 197 <212> PRT <213> Homo sapiens

<400> 1870 Met Arg Thr Leu Leu Thr Ile Leu Thr Val Gly Ser Leu Ala Ala His 10 Ala Pro Glu Asp Pro Ser Asp Leu Leu Gln His Val Lys Phe Gln Ser 20 25 Ser Asn Phe Glu Asn Ile Leu Thr Trp Asp Ser Gly Pro Glu Gly Thr 40 Pro Asp Thr Val Tyr Ser Ile Glu Tyr Lys Thr Tyr Gly Glu Arg Asp 55 60 Trp Val Ala Lys Lys Gly Cys Gln Arg Ile Thr Arg Lys Ser Cys Asn 70 75 Leu Thr Val Glu Thr Gly Asn Leu Thr Glu Leu Tyr Tyr Ala Arg Val 85 90 Thr Ala Val Ser Ala Gly Gly Arg Ser Ala Thr Lys Met Thr Asp Arg 105 Phe Ser Ser Leu Gln His Thr Thr Leu Lys Pro Pro Asp Val Thr Cys 120 Ile Ser Lys Val Arg Ser Ile Gln Met Ile Val His Pro Thr Pro Thr 135 140 Pro Ile Arg Ala Gly Asp Gly His Arg Leu Thr Leu Glu Asp Ile Phe 150 155 His Asp Leu Phe Tyr His Leu Glu Leu Gln Val Asn Arg Thr Tyr Gln 170 Met Val Ser Val Cys Cys Thr Leu Val Phe Leu Cys Leu Gly Ser Leu 185 Phe Pro Pro Asn 195 196

<210> 1871 <211> 75 <212> PRT <213> Homo sapiens

35 40 45

Arg Glu Ser Arg Ala Cys Ala Pro Gly Glu Arg Pro Asn Phe Leu Gly 50 55 60

Ile Arg Glu Gln Arg Leu Thr Gly Leu Val Val 65 70 75

<210> 1872 <211> 84 <212> PRT <213> Homo sapiens

<400> 1872 Met Pro Phe Ser Thr Cys Thr Ala Leu Pro Ser Trp Ala Thr Leu Ser 5 10 Thr Trp Ser Trp Thr Pro Lys Val Ser Leu Ala Gly Glu Glu Arg Gly 25 2.0 Glu Thr Cys Gln Pro Asp Pro Phe Pro Pro His Pro Ser Cys Ser Val 40 45 Gly Arg Thr Pro Pro His Ser Ser Leu Gly Ser Pro Pro Thr Thr Leu 55 60 Phe Leu Ser Pro Leu Leu Arg Val Glu Ser Arg Gly Ala Lys Cys Val 70 Val Cys Cys * 83

<210> 1873 <211> 51 <212> PRT <213> Homo sapiens

<210> 1874 <211> 503 <212> PRT <213> Homo sapiens

Glu	Trp		Leu	Gln	His	Asp		Ile	Pro	Gly	Asp	Leu 45	Arg	Asp	Leu
Arg	Val 50	35 Glu	Pro	Val	Thr	Thr 55	40 Ser	Val	Ala	Thr	Gly 60		Tyr	Ser	Ile
Leu 65		Asn	Val	ser	Trp 70	_	Leu	Arg	Ala	Asp 75	_	Ser	Ile	Arg	Leu 80
	Lys	Ala	Thr	Lys 85		Cys	Val	Thr	Gly 90		Ser	Asn	Phe	Gln 95	
Tyr	Ser	Cys	Val 100	Arg	Cys	Asn	Tyr	Thr 105		Ala	Phe	Gln	Thr 110	Gln	Thr
Arg	Pro	Ser 115	Gly	Gly	Lys	Trp	Thr 120	Phe	Ser	Tyr	Ile	Gly 125	Phe	Pro	Val
	130			Val		135					140				
145		•		Asp	150					155					160
_				His 165					170					175	
-			180	Asp				185					190		
		195		Asn			200					205			
	210			His		215					220				
225				Lys	230					235					240
	_			Gly 245					250					255	
			260	Cys				265					270		
		275		Pro			280	•				285			
_	290			Leu		295					300				
305				Ile	310					315					320
				Thr 325					330					335	
			340	Glu Asn				345					350		
		355					360					365			Thr
	370			Ala		375					380				
385				Asp	390					395					400
				405 Asp					410					415	
			420	Gln				425					430		
		435					440					445			
	450			Thr		455					460				Leu
465				Gln	470					475					480
				485			WTG	σту	490		SET	4111	AIG	495	114.0
Asp	сту	cys	cys	Ser	ьeu	^									

500 502

<210> 1875 <211> 158 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1) ... (158) <223> Xaa = any amino acid or nothing

<400> 1875 Met Xaa Pro Pro Thr Arg Pro Arg Thr Arg Gly Val Gly Ile Phe Tyr 10 5 Phe Val Ile Tyr Ile Ile Ser Phe Leu Val Val Asn Met Tyr 20 25 Ile Ala Val Ile Leu Glu Asn Phe Ser Val Ala Thr Glu Glu Ser Thr 40 Glu Pro Leu Ser Glu Asp Asp Phe Glu Met Phe Tyr Glu Val Trp Glu 55 Lys Phe Asp Pro Asp Ala Thr Gln Phe Ile Glu Phe Ser Lys Leu Ser 70 75 Asp Phe Ala Ala Leu Asp Pro Pro Leu Leu Ile Ala Lys Pro Asn 90 Lys Val Gln Leu Ile Ala Met Asp Leu Pro Met Val Ser Gly Asp Arg 105 Ile His Cys Leu Asp Ile Leu Phe Ala Phe Thr Lys Arg Val Leu Gly 120 125 Glu Ser Gly Glu Met Asp Ser Leu Arg Ser Gln Met Glu Glu Arg Phe 135 Met Ser Ala Asn Pro Ser Lys Val Ser Tyr Glu Pro Ile Thr 155

<210> 1876 <211> 106 <212> PRT <213> Homo sapiens

<400> 1876 Met Gly Asn Arg Ala Val Ile Ile Ala Arg Gln Leu Ser Ser Val His 1 5 10 Thr Leu Ile Cys Asn Phe Phe Trp Leu Leu Arg Thr Thr Gly Gly 25 Asp Leu Asp Ser Leu Lys Cys Ser Tyr Glu Ser Ile Gly Leu Asn Ser Ile Ser Thr His Glu Phe Ile Cys Thr Trp Gln Arg Arg Leu Asn Phe 60 55 Ser Phe Val Met Ser Phe Lys Pro Leu Phe Arg Ala Ser Pro His Ser 70 75 Tyr Leu Leu Ile Ile Gly Ser Gln Leu His Glu Thr Phe Asn Leu Gly 90 85 Ser Ile Ser Ser Glu Glu Lys Cys Ser 100 105

<210> 1877 <211> 241 <212> PRT

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<213> Homo sapiens
     <221> misc_feature
     <222> (1)...(241)
     <223> Xaa = any amino acid or nothing
     <400> 1877
Met Leu Trp Ala Leu Trp Pro Arg Trp Leu Ala Asp Lys Met Leu Pro
                                     10
                  5
Leu Leu Gly Ala Val Leu Leu Gln Lys Arg Glu Lys Arg Gly Pro Leu
                                 25
            20
Trp Arg His Trp Arg Arg Glu Thr Tyr Pro Tyr Tyr Asp Leu Gln Val
                             40
Lys Val Leu Arg Ala Thr Asn Ile Arg Gly Thr Asp Leu Leu Ser Lys
                         55
                                             60
Ala Asp Cys Tyr Val Gln Leu Trp Leu Pro Thr Ala Ser Pro Ser Pro
                     70
                                         75
Ala Gln Thr Arg Ile Val Ala Asn Cys Ser Asp Pro Glu Trp Asn Glu
                                     90
Thr Phe His Tyr Gln Ile His Gly Ala Val Lys Asn Val Leu Glu Leu
            100
                                105
Thr Leu Tyr Asp Lys Asp Ile Leu Gly Ser Asp Gln Leu Ser Leu Leu
                            120
Leu Phe Asp Leu Arg Ser Leu Lys Cys Gly Gln Pro His Lys His Thr
                        135
Phe Pro Leu Asn His Gln Asp Ser Gln Glu Leu Gln Val Glu Phe Val
                    150
                                        155
Leu Glu Lys Ser Gln Glu Pro Ala Ser Glu Val Ile Thr Asn Gly Val
                                    170
                165
Leu Gly Ala His Pro Trp Leu Arg Met Lys Gly Met Ile Leu Gly Glu
                                185
Gly Arg Ala Pro Arg Gln Gln His Gly Gln Ser Trp Glu Gly Gly Val
        195
                            200
Gly Pro Ser Pro Leu Ser Xaa Xaa Xaa Asn Thr Gly Gly Lys Ile Val
                                            220
                        215
Gly Phe Trp Glu Glu Met Ala Asn Gly Thr Gly Ala Pro Pro Arg Pro
225
                    230
                                        235
Pro
241
     <210> 1878
     <211> 50
     <212> PRT
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1028

10

Met Leu Leu Met Leu Leu Phe Arg Cys Cys Ser Ser Lys Asp Leu Trp

Pro Val Leu Ile Ala His Leu Val Pro Gln Gly Gly Gln Glu Gly Asn

<213> Homo sapiens

<400> 1878

25 Val Gly Glu Gln Thr Lys Gly Lys Ser Asn Arg Val Leu Pro Val Phe 40 Leu 49 <210> 1879 <211> 56 <212> PRT <213> Homo sapiens <400> 1879 Met Cys Ser Ala Phe Ser Ser Phe Trp Trp Val Pro Pro Leu Ala Gly 10 Ser Gly Val Lys Leu Gln Thr Phe Thr Ala Ser Val Thr Ala His Lys 20 25 Arg Ser Thr Asp Pro Lys Ser Glu Gln Gln Leu Asp Leu Ser Gln Arg 40 Thr Lys Glu Gln Ser Leu Thr Lys 55 56 <210> 1880 <211> 161 <212> PRT <213> Homo sapiens <221> misc feature <222> (1)...(161) <223> Xaa = any amino acid or nothing <400> 1880 Met Pro Ser Ala Ser Leu Leu Val Asn Leu Leu Ser Ala Leu Leu Ile 10 Leu Phe Val Phe Gly Glu Thr Glu Ile Arg Phe Thr Gly Gln Thr Glu 20 25 Phe Val Val Asn Glu Thr Ser Thr Thr Val Ile Arg Leu Ile Ile Glu Arg Ile Gly Glu Pro Ala Asn Val Thr Ala Ile Val Ser Leu Tyr Gly 55 Glu Asp Ala Gly Asp Phe Phe Asp Thr Tyr Ala Ala Ala Phe Ile Pro 75 Ala Gly Glu Thr Asn Arg Thr Val Tyr Ile Ala Val Cys Asp Asp Asp 90 Leu Pro Glu Pro Asp Glu Thr Phe Ile Phe His Leu Thr Leu Gln Lys 105 Pro Ser Ala Asn Val Lys Leu Gly Trp Pro Arg Thr Val Thr Val Thr 120 125 Ile Leu Ser Asn Gly Gln Met Ala Phe Trp Glu Phe Ile Phe Ile Leu 135 140

155

Asn Ile Gly Leu Pro Pro Pro Ile Pro Pro Ser Gly Xaa Leu Lys Ala

Pro 161

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<210> 1881
<211> 130
<212> PRT
<213> Homo sapiens
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<400> 1881 Met Gly Ile Tyr Gln Met Tyr Leu Cys Phe Leu Leu Ala Val Leu Leu 10 Gln Leu Tyr Val Ala Thr Glu Ala Ile Leu Ile Ala Leu Val Gly Ala 20 25 Thr Pro Ser Tyr His Trp Asp Leu Ala Glu Leu Leu Pro Asn Gln Ser 40 His Gly Asn Gln Ser Ala Gly Glu Asp Gln Ala Phe Gly Asp Trp Leu 55 Leu Thr Ala Asn Gly Ser Glu Ile His Lys His Val His Phe Ser Ser 70 Ser Phe Thr Ser Ile Ala Ser Glu Trp Phe Leu Ile Ala Asn Arg Ser 85 90 Tyr Lys Val Ser Ala Ala Ser Ser Phe Phe Ser Gly Val Phe Val 1.00 . 105 Gly Val Ile Ser Phe Gly Gln Leu Ser Asp Arg Phe Gly Arg Lys 120 Val Tyr 130

<210> 1882 <211> 108 <212> PRT <213> Homo sapiens

<400> 1882 Met Leu Trp Phe Ser Gly Val Gly Ala Leu Ala Glu Arg Tyr Cys Arg 10 Arg Ser Pro Gly Ile Thr Cys Cys Val Leu Leu Leu Asn Cys Ser 25 Gly Val Pro Met Ser Leu Ala Ser Ser Phe Leu Thr Gly Ser Val Ala 40 Lys Cys Glu Asn Glu Gly Glu Val Leu Gln Ile Pro Phe Ile Thr Asp 55 Asn Pro Cys Ile Met Cys Val Cys Leu Asn Lys Glu Val Thr Cys Lys 70 75 Arg Glu Lys Cys Pro Val Leu Ser Arg Asp Cys Ala Leu Ala Ile Lys 90 85 Gln Arg Gly Ala Cys Cys Glu Gln Cys Lys Gly Cys 108 100 105

<210> 1883 <211> 88 <212> PRT <213> Homo sapiens

<210> 1884 <211> 116 <212> PRT <213> Homo sapiens

<400> 1884 Met Cys Trp Ala Arg Cys Trp Thr Arg Trp Asn Thr Cys Thr Ile Trp Thr Ser Ser Thr Asp Pro Phe Arg Lys Cys Trp Met Ala Pro Glu Ala Leu Asn Phe Ser Phe Ser His Lys Ser Asp Ile Trp Ser Leu Gly Cys 40 Ile Ile Leu Asp Met Thr Ser Cys Ser Phe Met Asp Gly Thr Glu Ala 55 Met His Leu Arg Lys Ser Leu Arg Gln Ser Pro Gly Ser Leu Lys Ala 75 Val Leu Lys Thr Met Glu Glu Lys Gln Ile Pro Asp Val Glu Thr Phe 85 90 Arg Asn Leu Leu Pro Leu Met Leu Gln Ile Asp Pro Ser Asp Arg Ile 105 Thr Ile Lys * 115

<210> 1885 <211> 115 <212> PRT <213> Homo sapiens

<210> 1886 <211> 357 <212> PRT <213> Homo sapiens

<400> 1886 Met Ile Leu Ser Leu Leu Phe Ser Leu Gly Gly Pro Leu Gly Trp Gly 5 10 Leu Leu Gly Ala Trp Ala Gln Ala Ser Ser Thr Ser Leu Ser Asp Leu 25 Gln Ser Ser Arg Thr Pro Gly Val Trp Lys Ala Glu Ala Glu Asp Thr 40 Gly Lys Asp Pro Val Gly Arg Asn Trp Cys Pro Tyr Pro Met Ser Lys 55 Leu Val Thr Leu Leu Ala Leu Cys Lys Thr Glu Lys Phe Leu Ile His 70 Ser Gln Gln Pro Cys Pro Gln Gly Ala Pro Asp Cys Gln Lys Val Lys 85 90 Val Met Tyr Arg Met Ala His Lys Pro Val Tyr Gln Val Lys Gln Lys 105 Val Leu Thr Ser Leu Ala Trp Arg Cys Cys Pro Gly Tyr Thr Gly Pro 120 Asn Cys Glu His His Asp Ser Met Ala Ile Pro Glu Pro Ala Asp Pro 140 135 Gly Asp Ser His Gln Glu Pro Gln Asp Gly Pro Val Ser Phe Lys Pro 150 155 Gly His Leu Ala Ala Val Ile Asn Glu Val Glu Val Gln Gln Glu Gln 170 165 Gln Glu His Leu Leu Gly Asp Leu Gln Asn Asp Val His Arg Val Ala 185 Asp Ser Leu Pro Gly Leu Trp Lys Ala Leu Pro Gly Asn Leu Thr Ala 200 Ala Val Met Glu Ala Asn Gln Thr Gly His Glu Phe Pro Asp Arg Ser 215 220 Leu Glu Gln Val Leu Pro His Val Asp Thr Phe Leu Gln Val His 235 230 Phe Ser Pro Ile Trp Arg Ser Phe Asn Gln Ser Leu His Ser Leu Thr 250 . 245 Gln Ala Ile Arg Asn Leu Ser Leu Asp Val Glu Ala Asn Arg Gln Ala 260 265 Ile Ser Arg Val Gln Asp Ser Ala Val Ala Arg Ala Asp Phe Gln Glu 285 280 275 Leu Gly Ala Lys Phe Glu Ala Lys Val Gln Glu Asn Thr Gln Arg Val 295 300 Gly Gln Leu Arg Gln Asp Val Glu Asp Arg Leu His Ala Gln His Phe 310 315 Thr Leu His Arg Ser Ile Ser Glu Leu Gln Ala Asp Val Asp Thr Lys

325 330 Leu Lys Arg Leu His Lys Ala Gln Glu Ala Pro Gly Thr Asn Gly Ser 340 345 Leu Val Leu Glu Arg 355 357

<210> 1887 <211> 86 <212> PRT

<213> Homo sapiens

<400> 1887 Met Leu Cys Ser Arg Leu Gly Thr Thr Ala Ser Trp Arg Arg Leu Gly 10 Ile Arg Ala Trp Ala Pro Leu Leu Leu Phe Pro Trp Asp Trp His 25 Phe Ile Leu Ser Phe Ser Ser Arg Pro Trp Ala Gly Thr Leu Leu Ala 40 Pro His Asp Val Ile Met Gly Ser Ser Thr Phe Pro Gln Ser Cys Gln 55 Ala Glu Ala Gly Pro Arg His Ala Trp Pro Thr Gly Arg Phe Ser Arg Arg Leu Arg Arg Val

<210> 1888 <211> 48 <212> PRT <213> Homo sapiens

<400> 1888 Met Ser Val Arg Arg Ala Leu Thr Pro Ser Ala Leu Gly Leu Val Phe 5 10 Ile Leu Gln Ile Phe Ala His Gly Leu Pro Gly Pro Gly Pro Cys His 25 Leu Gly Pro Gly Ile Cys Leu Arg Ile Cys Gln Cys Ala Leu Asn * 40

<210> 1889 <211> 79 <212> PRT <213> Homo sapiens

<400> 1889 Met Ser Val Val Met Leu Ser Tyr Leu Leu Ser Ala Phe Phe Ser Gln 5 10 Ala Asn Thr Ala Ala Leu Cys Thr Ser Leu Val Tyr Met Ile Ser Phe 25 Leu Pro Tyr Ile Val Leu Leu Val Leu His Asn Gln Leu Ser Phe Val 40

Asn Gln Thr Phe Leu Cys Leu Leu Ser Thr Thr Ala Phe Gly Gln Gly 50 55 60

Val Phe Phe Ile Thr Phe Leu Glu Gly Gln Glu Thr Gly Ile His 65 70 75 79

<210> 1890 <211> 251 <212> PRT <213> Homo sapiens

<400> 1890 Met Asn Val Ile Tyr Phe Pro Leu His Leu Phe Val Val Tyr Ser Arg 10 Ala Tyr Thr Ser Leu Val Leu Val Gly Cys Thr Asn Leu Cys Ala Val 20 25 Leu Phe Ala Arg Cys Leu Asp Asp His Leu Val Ser Leu Arg Met Ser 40 45 Gly Ser Arg Lys Glu Phe Asp Val Lys Gln Ile Leu Lys Ile Arg Trp 60 55 Arg Trp Phe Gly His Gln Ala Ser Ser Pro Asn Ser Thr Val Asp Ser 70 75 Gln Gln Gly Glu Phe Trp Asn Arg Gly Gln Thr Gly Ala Asn Gly Gly 90 Arg Lys Phe Leu Asp Pro Cys Ser Leu Gln Leu Pro Leu Ala Ser Ile 105 Gly Tyr Arg Arg Ser Ser Gln Leu Asp Phe Gln Asn Ser Pro Ser Trp 120 Pro Met Ala Ser Thr Ser Glu Val Pro Ala Phe Glu Phe Thr Ala Glu 135 140 Asp Cys Gly Gly Ala His Trp Leu Asp Arg Pro Glu Val Asp Asp Gly 155 150 Thr Ser Glu Glu Glu Asn Glu Ser Asp Ser Ser Ser Cys Arg Thr Ser 170 165 Asn Ser Ser Gln Thr Leu Ser Ser Cys His Thr Met Glu Pro Cys Thr 185 Ser Asp Glu Phe Phe Gln Ala Leu Asn His Ala Glu Gln Thr Phe Lys 200 Lys Met Glu Asn Tyr Leu Arg His Lys Gln Leu Cys Asp Val Ile Leu 220 215 Val Ala Gly Asp Arg Arg Ile Pro Ala His Arg Leu Val Leu Ser Ser 230 235 Val Ser Asp Tyr Phe Ala Gly Met Phe Thr Asn

<211> 117
<212> PRT
<213> Homo sapiens

<221> misc_feature
<222> (1)...(117)
<223> Xaa = any amino acid or nothing

245

<210> 1891

<400> 1891 Met Leu Ile Asp Val Phe Phe Phe Leu Phe Leu Phe Ala Xaa Trp Met 10 Val Ala Phe Gly Val Ala Arg Gln Gly Ile Leu Arg Gln Asn Glu Gln 20 25 Arg Trp Arg Trp Ile Phe Arg Ser Val Ile Tyr Glu Pro Tyr Leu Ala 40 Met Phe Gly Gln Val Pro Ser Asp Val Asp Gly Thr Thr Tyr Asp Phe 55 Ala His Cys Thr Phe Thr Gly Asn Glu Ser Lys Pro Leu Cys Val Glu 75 Leu Asp Glu His Asn Leu Pro Arg Phe Pro Glu Trp Ile Thr Ile Pro 90 Leu Val Cys Ile Tyr Met Leu Ser Thr Asn Ile Leu Leu Val Asn Leu 100 105 Leu Val Ala Met Phe 115 117

<210> 1892 <211> 103 <212> PRT

<213> Homo sapiens

<400> 1892 Met Leu Cys His Pro His Val His His Leu Val Cys Leu Leu Ala Thr Leu Thr Phe Ser Leu Asn Ala Ser Cys Ala Glu Gln Thr Phe His 25 Ser Gln Gln Ser Asn Gly Glu Phe Met Ala Thr Leu Pro Ser Ile Ser 40 Lys Gln Phe Gly Val Ile Val Trp Lys Pro Gln Arg Lys Asp Val Ile 55 Arg Leu Pro Val Ala Leu Ser Phe Ser Ser Gly Ala Arg Leu Ala Phe 70 75 Thr Cys Leu Arg Lys Ile Ser Gly Phe Arg Ala Leu Ile Trp Gly Glu 85 90 Asp Lys Gly Trp Asp Leu * 100 102

<210> 1893 <211> 77 <212> PRT <213> Homo sapiens <221> misc_feature <222> (1)...(77) <223> Xaa = any amino acid or nothing

Ala Leu Val Phe Leu Leu Leu Val Gly Leu Leu Asn Ala Arg Gly Ile

35
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45

Lys Glu Ser Met Arg Ala Xaa Val Val Met Thr Val Val Glu Val Thr
50
55
60

Gly Leu Val Leu Val Val Val Leu Ala Leu Val Pro Gly
65
70
75
77

<210> 1894 <211> 46 <212> PRT <213> Homo sapiens

<210> 1895 <211> 162 <212> PRT <213> Homo sapiens

<400> 1895 Met Thr Ala Trp Arg Arg Phe Gln Ser Leu Leu Leu Leu Gly Leu . 10 5 Leu Val Leu Cys Ala Arg Leu Leu Thr Ala Ala Lys Gly Gln Asn Cys 25 20 Gly Gly Leu Val Gln Gly Pro Asn Gly Thr Ile Glu Ser Pro Gly Phe 40 35 Pro His Gly Tyr Pro Asn Tyr Ala Asn Cys Thr Trp Ile Ile Ile Thr 55 Gly Glu Arg Asn Arg Ile Gln Leu Ser Phe His Thr Phe Ala Leu Glu 70 Glu Asp Phe Asp Ile Leu Ser Val Tyr Asp Gly Gln Pro Gln Gln Gly 90 85 Asn Leu Lys Val Arg Leu Ser Gly Phe Gln Leu Pro Ser Ser Ile Val 105 Ser Thr Gly Ser Ile Leu Thr Leu Trp Phe Thr Thr Asp Phe Ala Val 125 120 Ser Ala Gln Gly Phe Lys Ala Leu Tyr Glu Gly Arg Arg Leu Val Val 140 135 Phe Cys Thr Cys Ile His Cys Pro Asn Asp Leu Ile His Ala Thr Leu 155 150 145 Asp * 161

<210> 1896 <211> 60

<212> PRT <213> Homo sapiens

<210> 1897 <211> 49 <212> PRT <213> Homo sapiens

<210> 1898 <211> 52 <212> PRT <213> Homo sapiens

<210> 1899 <211> 112 <212> PRT <213> Homo sapiens

<400> 1899

Met Ala Ile Pro Ser Val Val Ile Ser Gly Leu Ala Val Leu Leu Val 10 Ala Met Ala Leu Pro Ser Leu Ser Gly Ser Glu Ala Ile Lys Ser Met 25 Thr Ile Pro Gly Leu Val Val Pro Thr Val Val Arg Phe Met Ala Val Pro Gly Leu Ile Val Pro Ala Val Ala Lys Phe Thr Val Leu Pro Asp 55 Leu Thr Val Pro Thr Glu Asp Lys Ser Leu Ala Val Pro Ser Leu Ile 70 75 Ser Arg Ala Gly Asn Ser Val Pro Val Ser Ser Trp Asp Val Phe Gly 85 90 Val Ala Lys Leu Ile Ala Lys Leu Gly Leu Leu Ala Ala Ile Val Ala 100 105

<210> 1900

<211> 128

<212> PRT

<213> Homo sapiens

<400> 1900 Met Arg Val Tyr Gly Thr Cys Thr Leu Val Leu Met Ala Leu Val Val 10 Phe Val Gly Val Lys Tyr Val Asn Lys Leu Ala Leu Val Phe Leu Ala 25 20 Cys Val Val Leu Ser Ile Leu Ala Ile Tyr Ala Gly Val Ile Lys Ser 40 Ala Phe Asp Pro Pro Asp Ile Pro Val Cys Leu Leu Gly Asn Arg Thr 55 Leu Ser Arg Arg Ser Phe Asp Ala Cys Val Lys Ala Tyr Gly Ile His 75 70 Asn Asn Ser Ala Thr Ser Ala Leu Trp Gly Leu Phe Cys Asn Gly Ser 85 90 Gln Pro Ser Ala Ala Cys Asp Glu Tyr Phe Ile Gln Asn Asn Val Thr

100 105 110 Glu Ile Gln Gly Ile Pro Gly Ala Ala Ser Gly Val Phe Leu Glu Asn 115 120 125 128

<210> 1901

<211> 68

<212> PRT

<213> Homo sapiens

<400> 1901

Met Glu Leu Leu Lys Leu Leu Leu Thr Cys Phe Ser Glu Ala Met Tyr

1 5 10 15

Leu Pro Pro Ala Pro Glu Ser Gly Ser Thr Asn Pro Trp Val Gln Phe

20 25 30

Phe Cys Ser Thr Glu Asn Arg His Ala Leu Pro Leu Phe Thr Ser Leu

35 40 45

Leu Asn Thr Val Cys Ala Tyr Asp Pro Val Glu Tyr Gly Ile Pro Tyr
50 55 60

Asn His Leu Tyr
65 68

<210> 1902 <211> 127 <212> PRT <213> Homo sapiens

<400> 1902 Met Tyr Phe Ser Ser Leu Phe Pro Tyr Val Val Leu Ala Cys Phe Leu 1 5 10 Val Arg Gly Leu Leu Arg Gly Ala Val Asp Gly Ile Leu His Met 20 25 Phe Thr Pro Lys Leu Asp Lys Met Leu Asp Pro Gln Val Trp Arg Glu 40 Ala Ala Thr Gln Val Phe Ser Ala Leu Gly Leu Gly Phe Gly Gly Val 55 Ile Ala Phe Ser Ser Tyr Asn Lys Gln Asp Asn Asn Cys His Phe Asp 70 75 Ala Ala Leu Val Ser Phe Ile Asn Phe Phe Thr Ser Val Leu Ala Thr 85 90

Leu Val Val Phe Ala Val Leu Gly Phe Lys Ala Asn Ile Met Asn Glu
100 105 110

Lys Cys Val Val Glu Asn Ala Glu Lys Ile Leu Gly Tyr Arg Val
115 120 125 127

<210> 1903 <211> 83 <212> PRT <213> Homo sapiens

<400> 1903 Met Trp Lys Phe Val Ser Pro Leu Cys Met Ala Val Leu Thr Thr Ala 1 5 10 Ser Ile Ile Gln Leu Gly Val Thr Pro Pro Gly Tyr Ser Ala Trp Ile 20 25 Lys Glu Glu Ala Ala Glu Arg Tyr Leu Tyr Phe Pro Asn Trp Ala Met 40 Ala Pro Leu Ile Thr Leu Ile Val Val Ala Thr Leu Pro Ile Pro Val 55 Val Phe Val Leu Arg His Phe His Leu Ile Cys Asp Gly Ser Asn Thr 70 75 Pro Cys Ile 83

<210> 1904 <211> 129 <212> PRT

<213> Homo sapiens

<400> 1904 Met Lys Met Phe Val Ala His Gly Phe Tyr Ala Ala Lys Phe Val Val Ala Ile Gly Ser Val Ala Gly Leu Thr Val Ser Leu Leu Gly Ser Leu 25 Phe Pro Met Pro Arg Val Ile Tyr Ala Met Ala Gly Asp Gly Leu Leu 40 Phe Arg Phe Leu Ala His Val Ser Ser Tyr Thr Glu Thr Pro Val Val 55 60 Ala Cys Ile Val Ser Gly Phe Leu Ala Ala Leu Leu Ala Leu Leu Val 75 70 Ser Leu Arg Asp Leu Ile Glu Met Met Ser Ile Gly Thr Leu Leu Ala 90 85 Tyr Thr Leu Val Ser Val Cys Val Leu Leu Leu Arg His His Pro Glu 105 Ser Asp Ile Asp Gly Phe Val Lys Phe Leu Ser Glu Glu His Thr Cys 120 Ser 129

<210> 1905 <211> 93 <212> PRT <213> Homo sapiens

<400> 1905 Met Gly Leu Leu Met Met Ile Leu Gly Gln Ile Phe Leu Asn Gly Asn 10 Gln Ala Lys Glu Ala Glu Ile Trp Glu Met Leu Trp Arg Met Gly Val 25 20 Gln Arg Glu Arg Arg Leu Ser Ile Phe Gly Asn Pro Lys Arg Leu Leu 45 40 Ser Val Glu Phe Val Trp Gln Arg Tyr Leu Asp Tyr Arg Pro Val Thr 55 Asp Cys Lys Pro Val Glu Tyr Glu Phe Phe Trp Gly Pro Arg Ser His 70 Leu Glu Thr Thr Lys Met Lys Ile Leu Lys Phe Met Ala 85 90

<210> 1906 <211> 66 <212> PRT <213> Homo sapiens

40 Leu Ala Ser Gln His Ile Val Arg Thr Asp Leu His Val Gln Gly Pro Cys Ile 65 66

<210> 1907 <211> 105 <212> PRT

<213> Homo sapiens

<400> 1907 Met Leu Gln Leu Gly Pro Phe Leu Tyr Trp Thr Phe Leu Ala Ala Phe 10 Glu Gly Thr Val Phe Phe Gly Thr Tyr Phe Leu Phe Gln Thr Ala 25 Ser Leu Glu Glu Asn Gly Lys Val Tyr Gly Asn Trp Thr Phe Gly Thr 40 Ile Val Phe Thr Val Leu Val Phe Thr Val Thr Leu Lys Leu Ala Leu 55 60 Asp Thr Arg Phe Trp Thr Trp Ile Asn His Phe Val Ile Trp Gly Ser 70 75 Leu Ala Phe Tyr Val Phe Phe Ser Phe Phe Trp Gly Gly Ile Ile Trp 85 90 Pro Phe Leu Lys Gln Gln Arg Met Ala 100

<210> 1908 <211> 46 <212> PRT <213> Homo sapiens

<400> 1908 Met Gly Phe Leu Val Leu Lys Gln Pro Met Leu Val Ala Lys Val Phe 1 5 10 Pro Thr Leu Ala Gly Val Glu Ile Ile Leu Phe Thr Leu Lys Gly Phe 25 Pro Ile Leu Gly Ile Pro Val Gln Leu Pro Pro Thr Val *

<210> 1909 <211> 139 <212> PRT <213> Homo sapiens

<400> 1909 Met Ile Gln Ala Leu Gly Gly Phe Phe Thr Tyr Phe Val Ile Leu Ala 5 10 Glu Asn Gly Phe Leu Pro Ile His Leu Leu Gly Leu Arg Glu Asp Trp 25

<210> 1910

<211> 104

<212> PRT

<213> Homo sapiens

<400> 1910

Met Glu Gly Trp Phe Ala Val Leu Ser Thr Ala Asn Asp Val Leu Gly 10 Ala Pro Trp Asn Trp Leu Tyr Phe Ile Pro Leu Leu Ile Ile Gly Ala 20 25 Phe Phe Val Pro Thr Leu Val Leu Gly Val Leu Ser Gly Asp Phe Ala 40 Lys Glu Arg Glu Arg Val Glu Thr Arg Arg Ala Phe Met Lys Leu Arg 55 60 Arg Gln Gln Gln Ile Glu Arg Glu Leu Asn Gly Tyr Arg Val Trp Ile 75 70 Ala Lys Ala Glu Glu Val Met Leu Ala Glu Glu Asn Leu Tyr Pro Ser 85 His Ala Arg Pro Val Asn Pro * 100 103

<210> 1911

<211> 116

<212> PRT

<213> Homo sapiens

<400> 1911

 Met
 Ala
 Val
 Ala
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 Leu
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85 90 95

Pro Phe Ile Ser Arg Thr Lys Ile Ala Gln Leu Lys Ser Gly Arg Asp
100 105 110

Ser Thr Val *
115

<210> 1912 <211> 105 <212> PRT

<213> Homo sapiens

 A00> 1912

 Met Gln Leu Lys
 Thr Pro Ser Gly Gln Val Leu Ser Phe Cys Ile Leu 1
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 Gln Leu Phe Pro Phe Thr Ser Glu Ser Lys Arg Met Gly Val Ile Val 20
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<210> 1913 <211> 141 <212> PRT <213> Homo sapiens

<400> 1913 Met Leu Val Tyr Val Trp Ser Arg Arg Ser Pro Arg Val Arg Val Asn 1 5 10 Phe Phe Gly Leu Thr Phe Gln Ala Pro Phe Leu Pro Trp Ala Leu 25 Met Gly Phe Ser Leu Leu Gly Asn Ser Ile Leu Val Asp Leu Leu Gly Ile Ala Val Gly His Ile Tyr Tyr Phe Leu Glu Asp Val Phe Pro Asn Gln Pro Gly Arg Gln Glu Ala Pro Ala Asp Pro Trp Ala Phe Leu Lys Leu Leu Gly Cys Pro Cys Arg Arg Pro Gln Leu Thr Cys Pro 90 Ser Leu Arg Asn Ser Gln Asp Pro Ile Cys His Pro Arg Ser Ser Asp 105 , 110 Pro His Pro Gly Ala Arg Pro Lys Arg Leu Leu Ala Ala Ser Ile Leu 120 Pro Met Thr Pro Thr Trp Gly Arg Lys Asn Pro Ser * 135

<210> 1914 <211> 556 <212> PRT <213> Homo sapiens

<400> 1914 Met Lys Lys Val Leu Leu Leu Trp Lys Thr Val Leu Cys Thr Leu 5 10 Gly Gly Phe Glu Glu Leu Gln Ser Met Lys Ala Glu Lys Arg Ser Ile 25 Leu Gly Leu Pro Pro Leu Pro Glu Asp Ser Ile Lys Val Ile Arg Asn 40 Met Arg Ala Ala Ser Pro Pro Ala Ser Ala Ser Asp Leu Ile Glu Gln 55 Gln Gln Lys Arg Gly Arg Arg Glu His Lys Ala Leu Ile Lys Gln Asp 70 75 Asn Leu Asp Ala Phe Asn Glu Arg Asp Pro Tyr Lys Ala Asp Asp Ser 90 85 Arg Glu Glu Glu Glu Asn Asp Asp Asn Ser Leu Glu Gly Glu 100 105 Thr Phe Pro Leu Glu Arg Asp Glu Val Met Pro Pro Pro Leu Gln His 115 120 125 Pro Gln Thr Asp Arg Leu Thr Cys Pro Lys Gly Leu Pro Trp Ala Pro 135 140 Lys Val Arg Glu Lys Asp Ile Glu Met Phe Leu Glu Ser Ser Arg Ser 150 155 Lys Phe Ile Gly Tyr Thr Leu Gly Ser Asp Thr Asn Thr Val Val Gly 165 170 Leu Pro Arg Pro Ile His Glu Ser Ile Lys Thr Leu Lys Gln His Lys 185 Tyr Thr Ser Ile Ala Glu Val Gln Ala Gln Met Glu Glu Glu Tyr Leu 200 Arg Ser Pro Leu Ser Gly Gly Glu Glu Glu Val Glu Gln Val Pro Ala 215 Glu Thr Leu Tyr Gln Gly Leu Leu Pro Ser Leu Pro Gln Tyr Met Ile 230 235 Ala Leu Leu Lys Ile Leu Leu Ala Ala Pro Thr Ser Lys Ala Lys 245 250 Thr Asp Ser Ile Asn Ile Leu Ala Asp Val Leu Pro Glu Glu Met Pro 265 Thr Thr Val Leu Gln Ser Met Lys Leu Gly Val Asp Val Asn Arg His 280 Lys Glu Val Ile Val Lys Ala Ile Ser Ala Val Leu Leu Leu Leu 295 300 Lys His Phe Lys Leu Asn His Val Tyr Gln Phe Glu Tyr Met Ala Gln 310 315 His Leu Val Phe Ala Asn Cys Ile Pro Leu Ile Leu Lys Phe Phe Asn 325 330 Gln Asn Ile Met Ser Tyr Ile Thr Ala Lys Asn Ser Ile Ser Val Leu 345 Asp Tyr Pro His Cys Val Val His Glu Leu Pro Glu Leu Thr Ala Glu 360 Ser Leu Glu Ala Gly Asp Ser Asn Gln Phe Cys Trp Arg Asn Leu Phe 375 380 Ser Cys Ile Asn Leu Leu Arg Ile Leu Asn Lys Leu Thr Lys Trp Lys 390 395 His Ser Arg Thr Met Met Leu Val Val Phe Lys Ser Ala Pro Ile Leu

405 410 Lys Arg Ala Leu Lys Val Lys Gln Ala Met Met Gln Leu Tyr Val Leu 425 Lys Leu Leu Lys Val Gln Thr Lys Tyr Leu Gly Arg Gln Trp Arg Lys 440 Ser Asn Met Lys Thr Met Ser Ala Ile Tyr Gln Lys Val Arg His Arg 455 Leu Asn Asp Asp Trp Ala Tyr Gly Asn Asp Leu Asp Ala Arg Pro Trp 470 475 Asp Phe Gln Ala Glu Glu Cys Ala Leu Arg Ala Asn Ile Glu Arg Phe 485 490 Asn Ala Arg Arg Tyr Asp Arg Ala His Ser Asn Pro Asp Phe Leu Pro 505 Val Asp Asn Cys Leu Gln Ser Val Leu Gly Gln Arg Val Asp Leu Pro 525 520 Glu Asp Phe Gln Met Asn Tyr Asp Leu Trp Leu Glu Arg Glu Val Phe 535 Ser Lys Pro Ile Ser Trp Glu Glu Leu Leu Gln * 550

<210> 1915 <211> 212 <212> PRT

<213> Homo sapiens

<400> 1915 Met Phe Leu Val Ala Val Trp Trp Arg Phe Gly Ile Leu Ser Ile Cys 10 Met Leu Cys Val Gly Leu Val Leu Gly Phe Leu Ile Ser Ser Val Thr 20 25 Phe Phe Thr Pro Leu Gly Asn Leu Lys Ile Phe His Asp Asp Gly Val 40 Phe Trp Val Thr Phe Ser Cys Ile Ala Ile Leu Ile Pro Val Val Phe 55 Met Gly Cys Leu Arg Ile Leu Asn Ile Leu Thr Cys Gly Val Ile Gly 70 75 Ser Tyr Ser Val Val Leu Ala Ile Asp Ser Tyr Trp Ser Thr Ser Leu 90 85 Ser Tyr Ile Thr Leu Asn Val Leu Lys Arg Ala Leu Asn Lys Asp Phe 105 His Arg Ala Phe Thr Asn Val Pro Phe Gln Thr Asn Asp Phe Ile Ile 120 Leu Ala Val Trp Gly Met Leu Ala Val Ser Gly Ile Thr Leu Gln Ile Arg Arg Glu Arg Gly Arg Pro Phe Phe Pro Pro His Pro Tyr Lys Leu 150 155 Trp Lys Gln Glu Arg Glu Arg Arg Val Thr Asn Ile Leu Asp Pro Ser 170 Tyr His Ile Pro Pro Leu Arg Glu Arg Leu Tyr Gly Arg Leu Thr Gln 185 Ile Lys Gly Leu Phe Gln Lys Glu Gln Pro Ala Gly Glu Arg Thr Pro 195 200 Leu Leu Leu * 210 211

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<210> 1916
<211> 172
<212> PRT
<213> Homo sapiens
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<400> 1916 Met Cys Thr Pro Val Arg Val Ser Ile Val Cys Val Met Gly Ala Val 10 Gly Ala Val Trp Thr Ala Pro Leu Pro Leu Pro Trp Ala Pro Thr Pro 25 Ser Ile His Leu Arg Glu Glu Gly Ala Ala Phe Pro Phe Cys Gly Val 40 Cys Val Leu Arg Pro Arg Arg Ser Lys Trp Arg Ser Trp Asp Val Asn 55 Leu Gly Pro Arg Arg Gly Leu Leu Gly Cys Gly Pro Cys Pro Ser 70 Gly Lys Pro Arg Val His Leu Gln Arg Thr Arg Ser Gly Ala Gly Ala 85 90 Glu Ala Gly Gly Leu Pro Thr Arg Gly Ser Met Arg Gly Cys Pro Phe 105 100 Leu Gly Ser Ser Ala Ala Lys Cys Ser Leu Leu Arg Pro Pro Ser 120 125 Arg Gly Glu Ala Ser Pro Trp Leu Pro Glu Phe Met Thr His Pro Val 135 140 His His Gln Gln Leu Ala Cys Gly Ser Gly Trp Leu Gly Thr Lys His 150 155 Pro Gly Gly Thr Cys Ala Leu Gly Ser Thr Met * 165

<210> 1917 <211> 72 <212> PRT <213> Homo sapiens

<210> 1918 <211> 88 <212> PRT <213> Homo sapiens

<210> 1919 <211> 54 <212> PRT

<213> Homo sapiens

<210> 1920 <211> 114 <212> PRT <213> Homo sapiens

<400> 1920 Met His Pro Pro Leu Thr Pro Pro Thr Pro Leu Cys Leu Trp Leu Arg 1 5 10 Leu Leu Lys Ala Gln Ile Leu Ser Tyr Pro Val Pro Arg Phe Glu Thr 25 His Ser Leu Ile Ser Arg Cys Ser Gln Val Pro Pro Thr Phe Leu Trp Asp Ile Lys Lys Gly Val Arg Gly Gln Arg Glu Pro Ser Gly Pro Leu Leu Pro Tyr Thr Leu His Cys Pro Phe Ser Pro His Gln Asn Ala Gln 70 75 Arg Arg Cys Asp Asp Ala Thr Glu Asp Tyr Ala Thr Trp Ser Asn Arg 90 85 Ser Gly Gln His Asp Gln Leu Ser Arg Gly Cys Leu Leu Pro Phe Leu 105

Leu *

<210> 1921 <211> 139 <212> PRT <213> Homo sapiens

<400> 1921 Met Val Tyr Leu Tyr Ile Tyr Leu Asp Leu Phe Gln Phe Leu Ile Thr Val Leu Gln Gly Phe Leu Phe Val Phe Glu Met Glu Phe His Ser Cys 20 25 Arg Pro Gly Gln Ser Ala Met Met Gln Ser Gln Leu Ala Ala Thr Ser 40 45 Ala Ser Arg Val Gln Val Ile Leu Val Val Ser Ala Pro Gln Glu Ala 60 55 Gly Thr Thr Gly Ala Arg His His Val Gln Leu Ile Phe Val Phe Leu 70 75 Leu Glu Met Gly Phe Cys His Val Gly Gln Ala Gly Leu Glu Leu 85 90 Asn Ser Gly Asp Pro Pro Thr Ser Ala Ser Gln Ser Ala Gly Ile Arg 100 105 Gly Val Asn His Cys Ala Pro Pro Ile Asn Ser Leu Leu Thr Phe Gln 120 Ser Phe Ile His Leu Glu Cys Ile Val Ile *

135

<210> 1922 <211> 52 <212> PRT <213> Homo sapiens

<210> 1923 <211> 71 <212> PRT <213> Homo sapiens

35 40 45

Tyr Leu Leu Phe Phe Leu Trp Thr Phe Lys Leu Phe Ser Gly Phe Thr 50 55 60

Leu Lys Ile Ile Gln Gln *
65 70

<210> 1924 <211> 187 <212> PRT <213> Homo sapiens

<400> 1924 Met Leu Phe Ile Gln Tyr Leu Leu Pro Cys Leu Leu Leu Ser Ala Glu 10 Leu Ser Gly Thr Phe Phe Leu Tyr Asn Thr Cys His Leu His Val Pro 20 2.5 Cys Cys His Ser Leu Val Pro Thr Gly Pro Pro Ser Leu Ser Ser His 40 Phe Gln Ser Arg Gly Leu Cys Ala Pro Cys Ala Ser Ile Ala Asp Ser 55 Gly Ile Ala Asp Ser Gly Gly Asn Asn Leu Asn Phe Val Gly Ala Gly 70 75 Gly Val Ala Ser Gly His Leu Leu Ser Pro Leu Leu Gly Pro Gln Ser 85 90 Ser Pro Cys Pro His Cys Pro Arg Gly Gly Arg Leu Pro Ser Gln Pro 100 105 Leu Pro Leu Cys Ser Ala Arg Ser Trp Ala Gln Glu Ala Leu Arg Leu 120 125 Pro Ser Ser Ala Gln Leu Cys Pro Cys His Pro Leu Pro Arg Gly Leu 135 Gly Pro Val Ser Pro Ser Gly Leu Leu Ala Asn Ile Ser Tyr Arg His 155 Asn Trp Leu Leu Gly Ser Trp Pro Gly Trp Leu Ile Trp Gly Gly Lys 170 Asn Arg Gly Gly Leu Asn Ser Phe Leu Ala * 180 185 186

<210> 1925 <211> 50 <212> PRT <213> Homo sapiens

<210> 1926 <211> 47 <212> PRT <213> Homo sapiens

<210> 1927 <211> 149 <212> PRT <213> Homo sapiens

<400> 1927 Met Ala Thr Gly Leu Leu Ala Phe Leu Gly Leu Ala Ala Gly Gly Gln 10 Thr Leu Cys Pro Ala Gly Glu Leu Pro Gly His Ala Arg Ala Gln Ala 25 20 Ser Gly Ala Pro Gly Ser Val Leu Ile Ala Val Pro Gly Arg Arg 40 Val His Thr Cys Gly Pro Gly Pro Ala Ala Pro Ser Thr Arg Gly Glu 60 55 Cys Pro Pro Pro Ala Leu Gly His Thr Arg Pro Ala Arg Pro Arg Pro 75 70 Val Leu Leu Arg Pro Ser Cys Ser Pro Gly Ala Arg Gly Ala Gly Thr 90 85 Trp Cys Cys Ala Pro Ala Thr Gly His Ser Ala Pro Arg Gly Cys Pro 105 100 Pro Ala Arg Ala Ala Pro Thr Gly Ser Ala Thr Pro Ala Pro Pro 120 115 Ala Ala Cys Ala Ala Phe His Ser Ala Trp Ser Val Pro Pro Ala Gly 135 130 Arg Gln Gln Gly * 145 148

<210> 1928 <211> 446 <212> PRT <213> Homo sapiens

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40
Ile Ala Glu Cys Cys Ser Thr Pro Tyr Ser Leu Leu Gly Leu Val Phe
                        55
Thr Val Ser Phe Val Ala Leu Gly Val Leu Thr Leu Cys Lys Phe Tyr
                    70
                                       75
Leu Gln Gly Tyr Arg Ala Phe Met Asn Asp Pro Ala Met Asn Arg Gly
                85
                                   90
Met Thr Glu Gly Val Thr Leu Leu Ile Leu Ala Val Gln Thr Gly Leu
           100
                              105
Ile Glu Leu Gln Val Val His Arq Ala Phe Leu Leu Ser Ile Ile Leu
                          120
                                              125
Phe Ile Val Val Ala Ser Ile Leu Gln Ser Met Leu Glu Ile Ala Asp
                      135
                                         140
Pro Ile Val Leu Ala Leu Gly Ala Ser Arg Asp Lys Ser Leu Trp Lys
                  150
                                      155
His Phe Arg Ala Val Ser Leu Cys Leu Phe Leu Leu Val Phe Pro Ala
              165
                                  170
Tyr Met Ala Tyr Met Ile Cys Gln Phe Phe His Met Asp Phe Trp Leu
                              185
           180
Leu Ile Ile Ser Ser Ser Ile Leu Thr Ser Leu Gln Val Leu Gly
               200
                                             205
Thr Leu Phe Ile Tyr Val Leu Phe Met Val Glu Phe Arg Lys Glu
                   215
                                          220
Pro Val Glu Asn Met Asp Asp Val Ile Tyr Tyr Val Asn Gly Thr Tyr
       230
                                      235
Arg Leu Leu Glu Phe Leu Val Ala Leu Cys Val Val Ala Tyr Gly Val
              245
                                  250
Ser Glu Thr Ile Phe Gly Glu Trp Thr Val Met Gly Ser Met Ile Ile
                              265
Phe Ile His Ser Tyr Tyr Asn Val Trp Leu Arg Ala Gln Leu Gly Trp
                          280
Lys Ser Phe Leu Leu Arg Arg Asp Ala Val Asn Lys Ile Lys Ser Leu
                      295
Pro Ile Ala Thr Lys Glu Gln Leu Glu Lys His Asn Asp Ile Cys Ala
                                      315
Ile Cys Tyr Gln Asp Met Lys Ser Ala Val Ile Thr Pro Cys Ser His
               325
                                  330
Phe Phe His Ala Gly Cys Leu Lys Lys Trp Leu Tyr Val Gln Glu Thr
                              345
Cys Pro Leu Cys His Cys His Leu Lys Asn Ser Ser Gln Leu Pro Gly
                          360
Leu Gly Thr Glu Pro Val Leu Gln Pro His Ala Gly Ala Glu Gln Asn
                      375
                                          380
Val Met Phe Gln Glu Gly Thr Glu Pro Pro Gly Gln Glu His Thr Pro
                   390
                                      395
Gly Thr Arg Ile Gln Glu Gly Ser Arg Asp Asn Asn Glu Tyr Ile Ala
                                  410
               405
Arg Arg Pro Asp Asn Glu Glu Gly Ala Phe Asp Pro Lys Glu Tyr Pro
           420
                               425
His Ser Ala Lys Asp Glu Ala His Pro Val Glu Ser Ala *
                           440
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<210> 1929

<211> 120

<212> PRT

<213> Homo sapiens

<400> 1929 Met Val Leu Pro Leu Pro Trp Leu Ser Arg Tyr His Phe Leu Arg Leu 5 10 Leu Leu Pro Ser Trp Ser Leu Ala Pro Gln Gly Ser His Gly Cys Cys Ser Gln Asn Pro Lys Ala Ser Met Glu Glu Gln Thr Asn Ser Arg Gly 40 Asn Gly Lys Met Thr Ser Pro Pro Arg Gly Pro Gly Thr His Arg Thr Ala Glu Leu Ala Arg Ala Glu Glu Leu Glu Gln Gln Leu Glu Leu 75 Tyr Gln Ala Leu Leu Glu Gly Gln Glu Gly Ala Trp Glu Ala Gln Ala Leu Val Leu Lys Ile His Lys Leu Lys Glu Gln Met Arg Arg His Gln 100 105 Glu Ser Leu Gly Gly Gly Ala * 115

<210> 1930 <211> 122 <212> PRT <213> Homo sapiens

<400> 1930

Met Thr Trp Leu Val Leu Leu Gly Thr Leu Leu Cys Met Leu Arq Val 5 10 Gly Leu Gly Thr Pro Asp Ser Glu Gly Phe Pro Pro Arg Ala Leu His 20 25 Asn Cys Pro Tyr Lys Cys Ile Cys Ala Ala Asp Leu Leu Ser Cys Thr 40 Gly Leu Gly Leu Gln Asp Val Pro Ala Glu Leu Pro Ala Gly Thr Ala 55 Asp Leu Asp Leu Ser His Asn Ala Leu Gln Arg Met Arg Pro Gly Trp 70 Leu Ala Pro Leu Phe Gln Leu Arg Ala Leu His Leu Asp His Asn Glu 85 90 Leu His Ala Leu Asp Arg Gly Val Phe Val Asn Ala Ser Gly Leu Arg 105 Leu Leu Asp Leu Ser Ser Asn Ala Glu Phe 115 120

<210> 1931 <211> 73 <212> PRT <213> Homo sapiens

<400> 1931

 Met Ala Arg Ala Pro Ser Val Ala Leu Ala Gln Leu Trp Leu Ile Cys

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 10
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 Leu Cys Pro Glu Ser Leu Ala Ser Phe Val Gln Ala Val Pro Trp Lys
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 25
 30

 Val Leu Gln Pro Ser Ser Asn Arg Ser Thr Asp Cys Ser Pro His Met

35 40 45

Arg Pro Thr Cys Glu Thr Leu Gly Ser Arg Lys Ala Gln Asp Leu Gly 50 55 60

Ala Gly Tyr Tyr Val Ser Val His *
65 70 72

<210> 1932 <211> 68 <212> PRT <213> Homo sapiens

(213) Homo Baptens

<210> 1933 <211> 47 <212> PRT <213> Homo sapiens

<210> 1934 <211> 86 <212> PRT <213> Homo sapiens

Ala Val His Arg Lys Ala Gly Asp Thr Glu Val Gln Gln Ser Leu Leu 65 70 75 80

Leu Leu Leu Lys Lys *

<210> 1935

<211> 76

<212> PRT

<213> Homo sapiens

<400> 1935

<210> 1936

<211> 49

<212> PRT

<213> Homo sapiens

<400> 1936

 Met
 Leu
 Gln
 Thr
 Phe
 Val
 Thr
 Thr
 Cys
 Ile
 Ser
 Tyr
 Phe
 Tyr
 Trp

 1
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 His
 Phe
 Asn
 Phe
 Val
 Leu
 Ser

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 25
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 30
 30
 Phe
 Phe
 Gln
 Phe
 Leu
 Ile
 Gly
 Gln
 Val
 Tyr

 35
 40
 45
 48
 48

<210> 1937

<211> 76

<212> PRT

<213> Homo sapiens

<400> 1937

50 55 60
Glu Ile Lys Phe Tyr Ile Gln Leu Ala Lys Lys Lys
65 70 75 76

<210> 1938 <211> 191 <212> PRT <213> Homo sapiens

<400> 1938 Met Ala Asp Glu Lys Thr Phe Arg Ile Gly Phe Ile Val Leu Gly Leu 10 Phe Leu Leu Ala Leu Gly Thr Phe Leu Met Ser His Asp Arg Pro Gln 25 Val Tyr Gly Thr Phe Tyr Ala Met Gly Ser Val Met Val Ile Gly Gly 40 Ile Ile Trp Ser Met Cys Gln Cys Tyr Pro Lys Ile Thr Phe Val Pro 55 Ala Asp Ser Asp Phe Gln Gly Ile Leu Ser Pro Lys Ala Met Gly Leu 70 75 Leu Glu Asn Gly Leu Ala Ala Glu Met Lys Ser Pro Ser Pro Gln Pro 90 Pro Tyr Val Arg Leu Trp Glu Glu Ala Ala Tyr Asp Gln Ser Leu Pro 105 Asp Phe Ser His Ile Gln Met Lys Val Met Ser Tyr Ser Glu Asp His 120 Arg Ser Leu Leu Ala Pro Glu Met Gly Gln Pro Lys Leu Gly Thr Ser 135 Asp Gly Gly Glu Gly Pro Gly Asp Val Gln Ala Trp Met Glu Ala 150 155 Ala Val Val Ile His Lys Gly Leu Asn Glu Ser Glu Gly Glu Arg Arg

Leu Thr Gln Ser Trp Pro Gly Pro Leu Ala Cys Pro Gln Gly Pro

185

170

<210> 1939 <211> 82 <212> PRT <213> Homo sapiens

180

<210> 1940 <211> 101 <212> PRT <213> Homo sapiens

<400> 1940 Met His Val Cys Leu His Ile Trp Gly Leu Gly Val Cys Val Phe Met 10 His Met Met Cys Ala Cys Val Gly Val Tyr Val Cys Pro Phe Met Arg Tyr Gly Met Gln Ile Cys Ala Cys Ile His Ala His Ser Cys Ser Ala 40 Cys Val Cys Ser Cys Ile Trp Cys Met His Gly Cys Ser Tyr Leu Trp 55 Gly Thr Gly Ile Met His Val Cys Ser Ser Val Trp. Gly Val Gly Ile 75 70 Pro Gly Leu Trp Pro Glu Ala Pro Leu Gln Asp Thr Ala Pro Cys Arg 90 85 Leu Pro Arg Gly * 100

<210> 1941 <211> 88 <212> PRT <213> Homo sapiens

<210> 1942 <211> 46 <212> PRT <213> Homo sapiens

<400> 1942
Met Arg Ser Met Gly Phe Arg Ala Gln Gly Leu Pro Phe Gly Ile Arg
1 5 10 15
Gln Thr Trp Leu Arg Ile Leu Asp Leu Leu Thr Cys Thr Leu Pro

<210> 1943 <211> 155 <212> PRT <213> Homo sapiens

<400> 1943 Met Phe Thr Leu Leu Val Leu Leu Ser Gln Leu Pro Thr Val Thr Leu 10 Gly Phe Pro His Cys Ala Arg Gly Pro Lys Ala Ser Lys His Ala Gly 25 Glu Glu Val Phe Thr Ser Lys Glu Glu Ala Asn Phe Phe Ile His Arg 40 Arg Leu Leu Tyr Asn Arg Phe Asp Leu Glu Leu Phe Thr Pro Gly Asn 55 60 Leu Glu Arg Glu Cys Asn Glu Glu Leu Cys Asn Tyr Glu Glu Ala Arg 70 75 Glu Ile Phe Val Asp Glu Asp Lys Thr Ile Ala Phe Trp Gln Glu Tyr 85 90 Ser Ala Lys Gly Pro Thr Thr Lys Ser Asp Gly Asn Arg Glu Lys Ile 100 105 Asp Val Met Gly Leu Leu Thr Gly Leu Ile Ala Ala Gly Val Phe Leu 120 125 Val Ile Phe Gly Leu Leu Gly Tyr Tyr Leu Cys Ile Thr Lys Cys Asn 135 Arg Leu Gln His Pro Cys Ser Ser Ala Val Tyr

<210> 1944 <211> 61 <212> PRT <213> Homo sapiens

150

<210> 1945 <211> 79 <212> PRT <213> Homo sapiens | Met | Gln | Leu | Ile | Leu | Trp | Leu | Pro | Trp | Tyr | Val | Asp | Gln | Thr | Phe | Cys | Cys | Cys | Cys | Pro | Gly | Gln | Leu | Cys | Gln | Ser | Phe | Ser | Val | Leu | Gln | Cys | Cys | Cys | Pro | Gly | Gln | Leu | Cys | Gln | Ser | Phe | Ser | Asn | Arg | Asn | Asp | Ala | Arg | Leu | Leu | Gly | Ala | Lys | Gln | Ser | Ile | Ser | Asn | Arg | Arg | Trp | Leu | Glu | Pro | Ser | Val | Arg | Glu | Cys | Ala | Pro | Ser | Cys | Ser | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys | Cys

<210> 1946 <211> 72 <212> PRT <213> Homo sapiens

<210> 1947 <211> 56 <212> PRT <213> Homo sapiens

<210> 1948 <211> 48 <212> PRT <213> Homo sapiens

<400> 1948

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<210> 1949

<211> 136

<212> PRT

<213> Homo sapiens

<400> 1949

Met Leu Leu Ala Thr Leu Leu Leu Leu Leu Gly Gly Ala Leu Ala 10. His Pro Asp Arg Ile Ile Phe Pro Asn His Ala Cys Glu Asp Pro Pro 25 Ala Val Leu Leu Glu Val Gln Gly Thr Leu Gln Arg Pro Leu Val Arg 40 Asp Ser Arg Thr Ser Pro Ala Asn Cys Thr Trp Leu Ile Leu Gly Ser 55 60 Lys Glu Gln Thr Val Thr Ile Arg Phe Gln Lys Leu His Leu Ala Cys 70 75 Gly Ser Glu Arg Leu Thr Leu Arg Ser Pro Leu Gln Pro Leu Ile Ser 85 90 Leu Cys Glu Ala Pro Pro Ser Pro Leu Gln Leu Pro Gly Gly Asn Val 105 Thr Ile Thr Tyr Ser Tyr Ala Gly Ala Lys Arg Pro Gln Gly His Gly 120 Phe Phe Cys Phe Leu Lys Ala Lys 135 136

<210> 1950

<211> 78

<212> PRT

<213> Homo sapiens

<400> 1950

 Met Trp Ile Tyr Phe Trp Thr Leu Asn Ser Val Pro Val Ile Tyr Met

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 Ser Thr Leu Met Ser Ile Pro His Tyr Phe Asp Tyr Cys Cys Phe Ile
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 25
 30

 Val Ser Asp Ile Met Leu Pro Glu Ile Thr Phe Ser Thr Phe Ile Leu
 45

 Leu Leu Met Val Ala Leu Ala Ile Arg Gly Pro Leu His Phe Arg Arg
 50
 55
 60

 His Phe Arg Ile Asn Leu Ser Ile Ala Thr Lys Asn Ala *
 75
 77

<210> 1951

<211> 89 <212> PRT <213> Homo sapiens

<210> 1952 <211> 47 <212> PRT <213> Homo sapiens

<210> 1953 <211> 56 <212> PRT <213> Homo sapiens

<210> 1954 <211> 425 <212> PRT <213> Homo sapiens

<400> 1954 Met Thr Leu Arg Pro Gly Thr Met Arg Leu Ala Cys Met Phe Ser Ser 10 Ile Leu Leu Phe Gly Ala Ala Gly Leu Leu Phe Ile Ser Leu Gln 25 Asp Pro Thr Glu Leu Ala Pro Gln Gln Val Pro Gly Ile Lys Phe Asn 40 Ile Arg Pro Arg Gln Pro His His Asp Leu Pro Pro Gly Gly Ser Gln 55 Asp Gly Asp Leu Lys Glu Pro Thr Glu Arg Val Thr Arg Asp Leu Ser 75 Ser Gly Ala Pro Arg Gly Arg Asn Leu Pro Ala Pro Asp Gln Pro Gln 85 90 Pro Pro Leu Gln Arg Gly Thr Arg Leu Arg Leu Arg Gln Arg Arg Arg 100 105 Arg Leu Leu Ile Lys Lys Met Pro Ala Ala Thr Ile Pro Ala Asn 120 125 Ser Ser Asp Ala Pro Phe Ile Arg Pro Gly Pro Gly Thr Leu Asp Gly 135 140 Arg Trp Val Ser Leu His Arg Ser Gln Gln Glu Arg Lys Arg Val Met 150 155 Gln Glu Ala Cys Ala Lys Tyr Arg Ala Ser Ser Ser Arg Arg Ala Val 170 Thr Pro Arg His Val Ser Arg Ile Phe Val Glu Asp Arg His Arg Val 185 Leu Tyr Cys Glu Val Pro Lys Ala Gly Cys Ser Asn Trp Lys Arg Val 200 Leu Met Val Leu Ala Gly Leu Ala Ser Ser Thr Ala Asp Ile Gln His 215 Asn Thr Val His Tyr Gly Ser Ala Leu Lys Arg Leu Asp Thr Phe Asp 230 235 Arg Gln Gly Ile Leu His Arg Leu Ser Thr Tyr Thr Lys Met Leu Phe 245 250 Val Arg Glu Pro Phe Glu Arg Leu Val Ser Ala Phe Arg Asp Lys Phe 265 Glu His Pro Asn Ser Tyr Tyr His Pro Val Phe Gly Lys Ala Ile Leu 280 Ala Arg Tyr Arg Ala Asn Ala Ser Arg Glu Ala Leu Arg Thr Gly Ser 295 Gly Val Arg Phe Pro Glu Phe Val Gln Tyr Leu Leu Asp Val His Arg 310 315 Pro Val Gly Met Asp Ile His Trp Asp His Val Ser Arg Leu Cys Ser 325 330 Pro Cys Leu Ile Asp Tyr Asp Phe Val Gly Lys Phe Glu Ser Met Glu 345 Asp Asp Ala Asn Phe Phe Leu Ser Leu Ile Arg Ala Pro Arg Asn Leu Thr Phe Pro Arg Phe Lys Asp Arg His Ser Gln Glu Ala Arg Thr Thr 375 Ala Arg Ile Ala His Gln Tyr Phe Ala Gln Leu Ser Ala Leu Gln Arg 390 395 Gln Arg Thr Tyr Asp Phe Tyr Tyr Met Asp Tyr Leu Met Phe Asn Tyr 405 410 Ser Lys Pro Phe Ala Asp Leu Tyr * 420 424

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<210> 1955
<211> 106
<212> PRT
<213> Homo sapiens
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<400> 1955 Met Val Cys Phe Leu Phe Ile Thr Pro Leu Ala Ala Ile Ser Gly Trp 10 Leu Cys Leu Arg Gly Ala Gln Asp His Leu Arg Leu His Ser Gln Leu 25 Glu Ala Val Gly Leu Ile Ala Leu Thr Ile Ala Leu Phe Thr Ile Tyr 40 Val Leu Trp Thr Leu Val Ser Phe Arg Tyr His Cys Gln Leu Tyr Ser 55 Glu Trp Arg Lys Thr Asn Gln Lys Val Arg Leu Lys Ile Arg Glu Ala 70 75 Asp Ser Pro Glu Gly Pro Gln His Ser Pro Leu Ala Ala Gly Leu Leu 85 90 Lys Lys Val Ala Glu Glu Thr Pro Val *

<210> 1956 <211> 139 <212> PRT <213> Homo sapiens

100

<400> 1956 Met Val Leu Pro Phe Ile Cys Asn Leu Leu Arg Arg His Pro Ala Cys Arg Val Leu Val His Arg Pro His Gly Pro Glu Leu Asp Ala Asp Pro 25 Tyr Asp Pro Gly Glu Glu Asp Pro Ala Gln Ser Arg Ala Leu Glu Ser 40 Ser Leu Trp Glu Leu Gln Ala Leu Gln Arg His Tyr His Pro Glu Val 55 60 Ser Lys Ala Ala Ser Val Ile Asn Gln Ala Leu Ser Met Pro Glu Val 70 75 Ser Ile Ala Pro Leu Leu Glu Leu Thr Ala Tyr Glu Ile Phe Glu Arg 85 90 Asp Leu Lys Lys Lys Gly Pro Glu Pro Val Pro Thr Gly Val Leu Ser 100 105 Gln Pro Arg Ala Cys Trp Asp Gly Arg Val Lys Leu Cys Ala Gln His 115 120 Phe His Ala Gln Leu Thr Leu Ala His Leu * 130 135 138

<210> 1957 <211> 87 <212> PRT <213> Homo sapiens

<400> 1957 Met Ala Ala Pro Trp Arg Trp Pro Thr Gly Leu Leu Ala Val Leu 10 Arg Pro Leu Leu Thr Cys Arg Pro Leu Gln Gly Thr Thr Leu Gln Arg 25 Asp Gly Leu Leu Phe Glu His Asp Arg Gly Arg Phe Phe Thr Ile Leu 40 45 Gly Leu Val Cys Ala Gly Gln Gly Gly Phe Trp Ala Ser Met Ala Gly 55 60 Ala Gly Ala Leu Arg Thr Pro Gly Pro Leu Gln Gly Met Asn Val Glu 70 75 Arg His Glu Leu Leu Phe * 85 86

<210> 1958 <211> 48 <212> PRT

<213> Homo sapiens

<210> 1959 <211> 65 <212> PRT

<213> Homo sapiens

<210> 1960 <211> 78 <212> PRT <213> Homo sapiens

<400> 1960

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 Val
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 His
 Val
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 Ser
 Cys
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 Gly
 Gly
 Leu
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 Leu
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 Arg
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 Gly
 Gln
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 Ala
 Gln
 Arg
 Leu
 Gly
 His
 Cys

 Leu
 Trp
 Arg
 Ala
 Gly
 Glu
 Glu
 Leu
 Leu
 Pro
 Asn
 Ser
 Gly
 His

 Asp
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 Asp
 Gly
 Gly
 Pro
 Lys
 Asp
 Lys
 Gly
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<210> 1961 <211> 77

<212> PRT

<213> Homo sapiens

<400> 1961

 Met
 Trp
 Tyr
 Gly
 Val
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 Leu
 Trp
 Ala
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 Ser
 Ser
 Leu
 Phe
 Phe

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 His
 Val
 Pro
 Ala
 Gly
 Leu
 Ala
 Leu
 Phe
 Thr
 Leu
 Arg
 His
 His
 Lys
 Lys
 Gl
 Met
 Arg
 Thr
 Arg
 Ala
 Ala
 Ala
 Gly
 Leu
 Arg
 Thr
 His
 His
 Lys
 Glu
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 Ala
 Ala
 Ala
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 Arg
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 Thr
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 Arg
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 Leu
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<210> 1962 <211> 65

<212> PRT

<213> Homo sapiens

<400> 1962

 Met
 Phe
 Ser
 Ala
 Val
 Phe
 Pro
 Ala
 Val
 Ser
 Cys
 Gln
 Ile
 Ser
 Leu
 Leu

 Ser
 Thr
 Cys
 Asn
 Ser
 Leu
 Gln
 His
 Phe
 Pro
 Tyr
 Ala
 Gly
 Val
 Leu
 Cys

 Phe
 Arg
 Pro
 Val
 Leu
 Cys
 Leu
 Cys
 Pro
 Gly
 Gln
 Asp
 Phe
 Cys
 Gly
 Asn

 Val
 Arg
 Cys
 Gln
 Trp
 Arg
 Leu
 Leu
 Ala
 Gly
 Val
 Asp
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 Ser
 Asp
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 Gly
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<210> 1963

<211> 53

<212> PRT

<213> Homo sapiens

<221> misc_feature

<222> (1)...(53) <223> Xaa = any amino acid or nothing

<210> 1964 <211> 232 <212> PRT <213> Homo sapiens

<400> 1964 Met Pro Ser Val His Arg Leu Leu Gly Pro Gln Pro Val Pro Ser Arg 1 5 10 Arg Leu Arg Leu Ala Leu Leu Leu Ser Leu Gln Val Val Val 20 25 Phe Phe Leu Val Val Leu Gly Gln Gly Arg Leu Leu Gln Pro Cys Arg 40 Gly Cys Leu Glu Leu Pro Gly Gly Pro Gly Glu Ala Glu Asp His Gly Asp Leu Gly Gln Gly Trp Val Gly Leu Leu Gln Ala Leu Asp Pro Leu 70 Ser His Arg Arg Leu Val Met Ser Thr Arg His Ala His Gly Glu Asp 90 Arg Ala Phe Leu His Phe Ile Asp Val Lys Leu Val Val Val Pro Ala 100 105 Thr Pro His Ile Leu Gln Val Gln Leu His Arg Val Val Glu Val Pro 120 Leu Leu Arg Arg Leu Phe His Phe Pro Leu Leu Arg Gly Gln Gln Val 135 140 Ser Ser Glu Asp Val Val Ile His Thr Leu Val Ala Glu Pro Gln Gly 150 155 Glu Gly Ala Leu Asn Lys Asp Arg Pro Gly Trp Ile Val Ala Gly Gln 170 1.75 Gly Gly Leu Leu Ile Gly Thr Leu Asp Ser Trp Cys Gly Asp Ile His 180 185 Ala Leu Cys Pro Thr Met Trp Gly Trp Gly Gly Ser Ala Ala Pro Val 200 205 Glu Ser Leu Gly Lys Gly Thr Ser Gly Glu Gly Asp Gly Arg Arg Gln 215 220 Gly Gln Arg Thr Gly Pro Gly * 230 231

<210> 1965 <211> 253 <212> PRT

<213> Homo sapiens

<400> 1965 Met Gly Cys Ala Ile Ile Ala Gly Phe Leu His Tyr Leu Phe Leu Ala 10 Cys Phe Phe Trp Met Leu Val Glu Ala Val Ile Leu Phe Leu Met Val Arg Asn Leu Lys Val Val Asn Tyr Phe Ser Ser Arg Asn Ile Lys Met 40 Leu His Ile Cys Ala Phe Gly Tyr Gly Leu Pro Met Leu Val Val Val 55 Ile Ser Ala Ser Val Gln Pro Gln Gly Tyr Gly Met His Asn Arg Cys 70 75 Trp Leu Asn Thr Glu Thr Gly Phe Ile Trp Ser Phe Leu Gly Pro Val 85 90 Cys Thr Val Ile Val Ile Asn Ser Leu Leu Leu Thr Trp Thr Leu Trp 100 105 Ile Leu Arg Gln Arg Leu Ser Ser Val Asn Ala Glu Val Ser Thr Leu . 125 115 120 Lys Asp Thr Arg Leu Leu Thr Phe Lys Ala Phe Ala Gln Leu Phe Ile 135 Leu Gly Cys Ser Trp Val Leu Gly Ile Phe Gln Ile Gly Pro Val Ala 150 155 Gly Val Met Ala Tyr Leu Phe His His His Gln Gln Pro Ala Gly Gly 165 170 Leu His Leu Pro His Pro Leu Ser Ala Gln Arg Pro Gly Thr Arg Arg 185 Ile Gln Glu Val Asp His Trp Glu Asp Glu Ala Gln Leu Pro Val Pro 200 Asp Leu Lys Asp Leu Ala Val Leu His Ala Ile Arg Phe Gln Asp Gly 215 220 Leu Lys Ser Phe Leu Ala Phe Lys Tyr Ala Met Glu Pro Thr Val Gly 230 235 Gly Thr Ser Ser Phe Pro Cys Arg Glu Pro Tyr Pro 250

<210> 1966 <211> 649 <212> PRT <213> Homo sapiens

<400> 1966

 Met
 Val
 Thr
 Cys
 Phe
 Ile
 Ile
 Gly
 Leu
 Leu
 Phe
 Pro
 Val
 Phe
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 Val

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 Cys
 Tyr
 Leu
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 Met
 Ile
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Asn	Ser	Leu 115	Tyr	Leu	Ala	Thr	Ile 120	Ser	Leu	Lys	Ile	Val 125		Phe	Val
Lys	Tyr 130	Ser	Ala	Leu	Asn	Pro 135	Arg	Glu	Ser	Trp	Asp 140	Met	Trp	His	Pro
145			Ala		150					155					160
Leu	Arg	Leu	Ile	Ser 165	Leu	Phe	Thr	Ala	Asn 170	Ser	His	Leu	Gly	Pro 175	Leu
			Leu 180					185	_			_	190		
		195	Leu _				200				_	205			
	210		Tyr			215					220	_	_		
225			Gln		230					235					240
			Trp	245					250					255	
			Gln 260					265			_		270		
		275	Asn Asn	_			280					285			
	290		Phe			295					300			-	
305	110	د ړ د	1110	7114	310	1111	цуз	J.C.U	111	315	DCI	1 7 1	riie	Gru	320
Gly	Gly	Thr	Leu	Pro 325	Thr	Pro	Phe	Asn	Val 330	Ile	Pro	Ser	Pro	Lys 335	Ser
			Leu 340					345					350		
		355	Lys				360					365			
	370		Arg			375					380				
385			Tyr		390					395					400
			Glu	405					410					415	
			Glu 420 Ala					425					430		
		435	Ser				440					445			
	450		Phe			455					460				
465					470					475		****9		1120	480
Ile	Ala	Ser	Glu	Arg 485	His	Asn	Ile	Ser	Asn 490	Gly	Ser	Ala	Leu	Val 495	Val
		•	Pro 500					505	_				510		
		515	Phe				520					525			
	530		Ala			535					540				
545			Ala	_	550			_		555					560
στλ	ьeu	Ата	Ser	Arg 565	GTÀ	Asp	ьeu	ser	11e 570	Pro	дТÀ	ьeu	ser	G1u 575	GIN

 Cys
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 Leu
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 Asp
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 Asp
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 Gly
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 Gly
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 Gly
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 Gly
 Leu
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<210> 1967 <211> 80 <212> PRT

<213> Homo sapiens

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<210> 1968 <211> 49 <212> PRT <213> Homo sapiens

<210> 1969 <211> 150 <212> PRT <213> Homo sapiens

<400> 1969
Met His Val His Phe Trp Leu Val Thr Ala Ser Phe Ser Ser Val

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<210> 1970

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1970

<210> 1971

<211> 64

<212> PRT

<213> Homo sapiens

<400> 1971

 Met Leu Ile Phe Thr Val Leu Glu Leu Leu Leu Ala Ala Tyr Ser Ser 1
 5
 10
 15

 Val Phe Trp Trp Lys Gln Leu Tyr Ser Asn Asn Pro Gly Val Ser Met 20
 25
 30

 Leu Thr Cys Arg Leu Ile Pro Ala Val Ser Gln Val Gln Ala Thr Ile 35
 40
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 Ile Gln Pro Gln Lys Val Ala Lys Arg Arg Ile Asn Tyr Cys Ser *
 50
 63

<210> 1972

<211> 211

<212> PRT

<213> Homo sapiens <221> misc_feature <222> (1)...(211) <223> Xaa = any amino acid or nothing <400> 1972 Met Thr Arg Met Leu Asn Met Leu Ile Val Phe Arg Phe Leu Arg Ile 5 10 Ile Pro Ser Met Lys Pro Met Ala Val Val Ala Ser Thr Val Leu Gly 25 Leu Val Gln Asn Met Arg Ala Phe Gly Gly Ile Leu Val Val Val Tyr 40 Tyr Val Phe Ala Ile Ile Gly Ile Asn Leu Phe Arg Gly Val Ile Val 55 Ala Leu Pro Gly Asn Ser Ser Leu Ala Pro Ala Asn Gly Ser Ala Pro 75 70 Cys Gly Ser Phe Glu Gln Leu Glu Tyr Trp Ala Asn Asn Phe Asp Asp 90 85 Phe Xaa Ala Ala Leu Val Thr Leu Trp Asn Leu Met Val Val Asn Asn 100 105 Trp Gln Val Phe Leu Asp Ala Tyr Arg Arg Tyr Ser Gly Pro Trp Ser 125 120 Lys Ile Tyr Phe Val Leu Trp Trp Leu Val Ser Ser Val Ile Trp Val 135 140 Asn Leu Phe Leu Ala Leu Ile Leu Glu Asn Phe Leu His Lys Trp Asp 155 150 Pro Arg Ser His Leu Gln Pro Leu Ala Gly Thr Pro Glu Ala Thr Tyr 170 Gln Met Thr Val Glu Leu Leu Phe Arg Asp Ile Leu Glu Glu Pro Gly 185 Glu Asp Glu Leu Thr Glu Arg Leu Ser Gln His Pro His Leu Trp Leu 195 Cys Arg 210 <210> 1973 <211> 53 <212> PRT <213> Homo sapiens <400> 1973 Met Ile Gln Tyr Ala Val Phe Val Leu Cys Gly Phe Leu Tyr Leu Cys 5 10 Phe Met Leu Phe Phe Ser Ser Val Thr Gln Ala Gly Val Ser Glu 25 Pro Arg Ser Ser His Cys Thr Pro Ala Trp Ala Thr Glu Arg Asp Cys 35 Val Ser Asn Lys 50 52

<210> 1974 <211> 50

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<210> 1975 <211> 87 <212> PRT <213> Homo sapiens

<210> 1976 <211> 107 <212> PRT <213> Homo sapiens

<210> 1977 <211> 134 <212> PRT <213> Homo sapiens

<400> 1977 Met Val Thr Val Ala Met Ala Cys Ser Gly Ala Leu Thr Ala Leu Cys -5 10 Cys Leu Phe Val Ala Met Gly Val Leu Arg Val Pro Trp His Cys Pro 25 Leu Leu Leu Val Thr Glu Gly Leu Leu Asp Met Leu Ile Ala Gly Gly Tyr Ile Pro Ala Leu Tyr Phe Tyr Phe His Tyr Leu Ser Ala Ala Tyr 55 Gly Ser Pro Val Cys Lys Glu Arg Gln Ala Leu Tyr Gln Ser Lys Gly 70 Tyr Ser Gly Phe Gly Cys Ser Phe His Gly Ala Asp Ile Gly Ala Gly 90 85 Ile Phe Ala Ala Leu Gly Ile Val Val Phe Ala Leu Gly Ala Val Leu 105 Ala Ile Lys Gly Tyr Arg Lys Val Arg Lys Leu Lys Glu Lys Pro Ala 120 115 Glu Met Phe Glu Phe * 133 130

<210> 1978 <211> 61 <212> PRT <213> Homo sapiens

<210> 1979 <211> 66 <212> PRT <213> Homo sapiens

20 25 30

Arg His Lys Tyr Ser Tyr Asp Ala Asn Val Phe Leu Gln Val Asn Tyr 35 40 45

Ile Thr Trp Pro Asp Ser Phe Ser Pro Val Pro Ser Leu Pro Pro Ile 50 55 60

Leu *
65

<210> 1980 <211> 51 <212> PRT <213> Homo sapiens

<210> 1981 <211> 79 <212> PRT <213> Homo sapiens

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<210> 1982 <211> 156 <212> PRT <213> Homo sapiens

Asn Tyr Asp Ile Cys Lys Val Tyr Leu Ala Arg Trp Gly Ile Gln Gly 40 Arg Trp Met Lys Gln Asp Pro Arg Arg Trp Gly Asn Pro Ala Arg Ala 55 Pro Arg Pro Gly Gln Arg Ala Pro Gln Pro Gln Pro Pro Pro Gly Pro Leu Pro Gln Ala Pro Gln Ala Val His Thr Leu Arg Gly Asp Ala His 90 Ser Pro Pro Leu Met Thr Phe Gln Ser Ser Ser Ala Trp Glu Gly Ala 100 105 Ser Gln Gln Glu Ile Pro Glu Asn Glu Glu Thr Glu Lys Gly Asp 120 115 125 Asp Gln Ile Ser Ser Phe Leu Gly Val Thr Ser Asn Thr Lys Glu Ala 135 140 Ser Val Ile Gly Ile Gln Lys Thr Val Asp Val Leu 150

<210> 1983 <211> 63 <212> PRT

<213> Homo sapiens

<210> 1984 <211> 232 <212> PRT <213> Homo sapiens

<400> 1984 Met Phe His Arg Cys Gly Ile Met Ala Leu Val Ala Ala Tyr Leu Asn 10 Phe Val Ser Gln Met Ile Ala Val Pro Ala Phe Cys Gln His Val Ser 25 Lys Val Ile Glu Ile Arg Thr Met Glu Ala Pro Tyr Phe Leu Pro Glu 40 His Ile Phe Arg Asp Lys Cys Met Leu Pro Lys Ser Leu Glu Lys His 55 Glu Lys Asp Leu Tyr Phe Leu Thr Asn Lys Ile Ala Glu Ser Leu Gly 70 75 Gly Lys Trp Asp Ile Val Leu Arg Asp Cys Gln Phe Arg Met Leu Pro 85 90 Gln Val Thr Asp Glu Asp Arg Leu Ser Arg Arg Lys Ser Ile Val Asp 105 Thr Val Ser Ile Gln Val Asp Ile Leu Ser Asn Asn Val Pro Ser Asp

115 120 125 Asp Val Val Ser Asn Thr Glu Glu Ile Thr Phe Glu Ala Leu Lys Lys 135 Ala Ile Asp Thr Ser Gly Met Glu Glu Glu Lys Glu Lys Arg Arg 150 155 Leu Val Ile Glu Lys Phe Gln Lys Ala Pro Phe Glu Glu Ile Ala Ala 165 170 Gln Cys Glu Ser Lys Ala Asn Leu Leu His Asp Arg Leu Ala Gln Ile 185 180 Leu Glu Leu Thr Ile Arg Pro Pro Pro Ser Pro Ser Gly Thr Leu Thr 200 205 Ile Thr Ser Gly His Ala Gln Tyr Gln Ser Val Pro Val Tyr Glu Met 215 220 Lys Phe Pro Asp Leu Cys Val Tyr 230 232

<210> 1985 <211> 141 <212> PRT <213> Homo sapiens

<400> 1985 Met Asn Leu Ser Leu Pro Phe Leu Cys Leu Phe Leu Ser Phe Ser 10 5 Phe Lys Leu Ala Leu Gln Leu Arg Lys Val Ser Leu Leu Ser Leu Arg 20 25 Leu Trp Gly Gln Ser Ile Cys Cys Leu Glu Lys Glu Gly Asn Gln Asp 40 Ser Ser Gly Thr Gln Met Ser Ser Leu Ala Leu Leu Asn Pro Leu 55 Leu His Asn Trp Ser Phe Ile Leu Ala Leu Asn Asp Pro Ala Gly His 75 70 His Gly Phe Leu Phe Leu Leu Val Phe Phe Ser Glu Thr Glu Ser 90 85 His Ser Val Thr Gln Ala Gly Val Gln Trp Arg Asp Leu Ser Ser Leu 100 105 Gln Pro Leu Pro Pro Gly Phe Lys Arg Phe Phe Cys Leu Ser Leu Pro

120

Ser Ser Trp Asp Tyr Arg Cys Ala Thr Thr Pro Gly *

<210> 1986 <211> 292 <212> PRT <213> Homo sapiens

Asn Glu Thr Leu Lys His Leu Thr Asn Asp Thr Thr Thr Pro Glu Ser 55 Thr Met Thr Ser Gly Gln Ala Arg Ala Ser Thr Gln Ser Pro Gln Ala 7.5 Leu Glu Asp Ser Gly Pro Val Asn Ile Ser Val Ser Ile Thr Leu Thr 90 Leu Asp Pro Leu Lys Pro Phe Gly Gly Tyr Ser Arg Asn Val Thr His 100 105 Leu Tyr Ser Thr Ile Leu Gly His Gln Ile Gly Leu Ser Gly Arg Glu 120 125 Ala His Glu Glu Ile Asn Ile Thr Phe Thr Leu Pro Thr Ala Trp Ser 135 140 Ser Asp Asp Cys Ala Leu His Gly His Cys Glu Gln Val Val Phe Thr 150 155 Ala Cys Met Thr Leu Thr Ala Ser Pro Gly Val Phe Pro Val Thr Val . 165 170 Gln Pro Pro His Cys Val Pro Asp Thr Tyr Ser Asn Ala Thr Leu Trp 180 185 Tyr Lys Ile Phe Thr Thr Ala Arg Asp Ala Asn Thr Lys Tyr Ala Gln 195 200 205 Asp Tyr Asn Pro Phe Trp Cys Tyr Lys Gly Ala Ile Gly Lys Val Tyr 215 220 His Ala Leu Asn Pro Lys Leu Thr Val Ile Val Pro Asp Asp Asp Arg 230 235 Ser Leu Ile Asn Leu His Leu Met His Thr Ser Tyr Phe Leu Phe Val 245 250 Met Val Ile Thr Met Phe Cys Tyr Ala Val Ile Lys Gly Arg Pro Ser 260 265 Lys Leu Arg Gln Ser Asn Pro Glu Phe Cys Pro Glu Lys Val Ala Leu 275 280 Ala Glu Ala * 290 291

<210> 1987 <211> 186 <212> PRT <213> Homo sapiens

<400> 1987 Met Ala Gly Pro Arg Pro Arg Trp Arg Asp Gln Leu Leu Phe Met Ser 10 Ile Ile Val Leu Val Ile Val Ile Cys Leu Met Leu Tyr Ala Leu 25 Leu Trp Glu Ala Gly Asn Leu Thr Asp Leu Pro Asn Leu Arg Ile Gly 40 Phe Tyr Asn Phe Cys Leu Trp Asn Glu Asp Thr Ser Thr Leu Gln Cys 55 His Gln Phe Pro Glu Leu Glu Ala Leu Gly Val Pro Arg Val Gly Leu 70 75 . Gly Leu Ala Arg Leu Gly Val Tyr Gly Ser Leu Val Leu Thr Leu Phe 85 90 Ala Pro Gln Pro Leu Leu Ala Gln Cys Asn Ser Asp Glu Arg Ala 105 Trp Arg Leu Ala Val Gly Phe Leu Ala Val Ser Ser Val Leu Leu Ala Gly Gly Leu Gly Leu Phe Leu Ser Tyr Val Trp Lys Trp Val Arg Leu

<210> 1988 <211> 47 <212> PRT <213> Homo sapiens

<210> 1989 <211> 58 <212> PRT <213> Homo sapiens

<210> 1990 <211> 80 <212> PRT <213> Homo sapiens

Thr His Trp Ala Val Cys Gly Cys Gly Phe Ile Ser Glu Lys Leu * 65 70 75 79

<210> 1991

<211> 48

<212> PRT

<213> Homo sapiens

<400> 1991

<210> 1992

<211> 51

<212> PRT

<213> Homo sapiens

<400> 1992

 Met Leu Phe Ser Leu Gln Thr Ala Ile Val Tyr Cys Thr Ile Thr Val

 1
 5
 10
 15

 Leu Cys His Arg Thr Leu Ile Phe Ser Ser Met His Lys Cys Ile Met
 20
 25
 30

 Leu Phe Pro Ile Ile His Ile Cys Ser Tyr Val Phe Phe Val Ile Tyr
 35
 40
 45

 Ser Phe *
 50

<210> 1993

<211> 79

<212> PRT

<213> Homo sapiens

<400> 1993

 Met
 Trp
 Cys
 Ala
 Glu
 Met
 Leu
 His
 Ile
 Leu
 Phe
 Met
 Gly
 Leu
 Arg
 Val

 Asn
 Leu
 Asn
 His
 Glu
 Thr
 Phe
 Leu
 Ile
 Ile
 Cys
 Cys
 Glu
 Ile
 Tyr
 Gln

 Ala
 Trp
 Met
 Ile
 Ser
 Val
 Phe
 Leu
 Val
 Val
 Cys
 Cys
 Phe
 Phe
 Lys
 Glu

 Val
 Ile
 Gln
 Val
 Pro
 Leu
 Leu
 Ser
 Cys
 Gln
 His
 Thr
 Lys
 Leu
 Lys

 Val
 Ile
 Gln
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 Fro
 Fro
 Arg
 Ser
 Arg
 Ser
 Gln
 Pro
 Val
 Glu
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 Lys
 Leu
 Thr
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 Ser
 Phe
 Arg
 Ser
 Arg
 Ser
 Gln
 Pro
 Val
 Glu
 *

 Lys
 Leu
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 Ile
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 Arg
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<210> 1994 <211> 52 <212> PRT <213> Homo sapiens

<210> 1995 <211> 164 <212> PRT <213> Homo sapiens

<400> 1995 Met Leu Leu Ala Thr Leu Leu Leu Leu Leu Gly Gly Ala Leu Ala . His Pro Asp Arg Ile Ile Phe Pro Asn His Ala Cys Glu Asp Pro Pro 25 Ala Val Leu Leu Glu Val Gln Gly Thr Leu Gln Arg Pro Leu Val Arg 40 Asp Ser Arg Thr Ser Pro Ala Asn Cys Thr Trp Leu Ile Leu Gly Ser 55 Lys Glu Arg Thr Val Thr Ile Arg Phe Gln Lys Leu His Leu Ala Cys 70 75 Gly Ser Glu Arg Leu Thr Leu Arg Ser Pro Leu Gln Pro Leu Ile Ser 85 90 Leu Cys Glu Ala Pro Pro Ser Pro Leu Gln Leu Pro Gly Gly Asn Val 105 Thr Ile Thr Tyr Ser Tyr Ala Gly Gly Gln Ser Thr His Gly Pro Gly 120 Leu Pro Ala Leu Leu Gln Ala Ser Pro Ser Pro Trp Cys Leu Cys Arg 135 140 Leu Ala Asp Val Leu Ala Arg Arg Gly Ser Met Pro Glu Pro Pro Leu 155 Cys Ile Cys * 163

<210> 1996 <211> 77 <212> PRT <213> Homo sapiens

His Val Pro Ala Gly Leu Leu Ala Leu Phe Thr Leu Arg His His Lys 20 25 30

Tyr Gly Ala Ala Ile Ala Gly Val Tyr Arg Ala Ala Gly Lys Glu Met 35 40 45

Ile Pro Phe Glu Ala Leu Thr Leu Gly Thr Gly Gln Thr Phe Cys Val 50 55 60

Leu Val Val Ser Phe Leu Arg Ile Leu Ala Thr Leu * 75 76

<210> 1997 <211> 233 <212> PRT <213> Homo sapiens

<400> 1997 Met Gly Leu Pro Gly Leu Phe Cys Leu Ala Val Leu Ala Ala Ser Ser 10 Phe Ser Lys Ala Arg Glu Glu Glu Ile Thr Pro Val Val Ser Ile Ala 20 25 Tyr Lys Val Leu Glu Val Phe Pro Lys Gly Arg Trp Val Leu Ile Thr 40 Cys Cys Ala Pro Gln Pro Pro Pro Ile Thr Tyr Ser Leu Cys Gly 55 60 Thr Lys Asn Ile Lys Val Ala Lys Lys Val Val Lys Thr His Glu Pro 70 75 Ala Ser Phe Asn Leu Asn Val Thr Leu Lys Ser Ser Pro Asp Leu Leu 90 Thr Tyr Phe Cys Arg Ala Ser Ser Thr Ser Gly Ala His Val Asp Ser 105 Ala Arg Leu Gln Met His Trp Glu Leu Trp Ser Arg Gln Arg Gly Arg 125 120 Pro Gln Gly Gly Asp Asp Leu Pro Gly Val Leu Gly Gln Pro Thr Tyr 135 His Gln Gln Pro Asp Arg Glu Gly Trp Ala Gly Pro Pro Ala Ala Glu 155 150 Thr Met Pro Gln Glu Ala Cys Gln Leu Ser Pro Ser Cys Arg Ala Arg 170 165 His Arg Thr Trp Phe Trp Cys Gln Ala Cys Lys Gln Arg Gln Cys Ser 185 180 Ser Thr Ala Pro Ser Gln Trp Leu Pro Gln Val Val Thr Gln Lys Met 200 205 Glu Asp Trp Gln Gly Pro Pro Gly Glu Pro His Pro Cys Leu Ala Ala 215 Leu Gln Glu His Pro Pro Ser Glu * 225 230 232

<210> 1998 <211> 58 <212> PRT <213> Homo sapiens

<400> 1998
Met Pro Ala Ile Val Val Phe Leu Phe Cys Phe Val Ile Ser Asp Gly

5 10 Leu Thr Leu Ser Pro Arg Leu Asp Cys Thr Gly Leu Asn Leu Leu Ser 20 25 3.0 Ser Ser Asp Arg Pro Thr Ser Ala Ser Pro Val Ala Gly Thr Ile Ala 40 Val Gln His His Ala Trp Leu Ile Phe 55

<210> 1999 <211> 66 <212> PRT <213> Homo sapiens

<400> 1999 Met Trp Leu Leu Val Thr Leu Ser Pro Arg Leu Leu Ser Pro Ser 10 His Phe Thr Leu Glu Gly Pro Gln Ile Asp Gln Ala His Ser Glu Leu 25 Gln Val Leu Pro Leu Val Arg Pro Ser Ala Val Pro Leu Leu Gln Arg 40 45 Ala Ser Trp Leu Arg Ser Arg Cys Leu His Leu Pro Lys Thr Val Leu 55 Val 65

<210> 2000 <211> 106 <212> PRT <213> Homo sapiens

<400> 2000 Met Gly Arg Cys Leu Ser Leu Gly Ile Leu Arg Gln Gly Leu Cys Cys 5 Pro Cys Trp Ser Val Val Ala Glu Ser Gly Leu Thr Ala Ser Leu Gly 20 Gly Ser Gly His Pro Ala Thr Ser Cys Ser Lys Glu Ala Gly Thr Thr 40 Gly Glu Cys Met His His Thr Gln Leu Gly Ile Gln Thr Leu Arg Thr 55 Tyr Tyr Met Pro Asp Ser Val Glu Leu Ser Glu Thr Met Ser Gly Cys 70 75 Asn Trp Leu Pro Thr Gln Gln Thr Gln Ser Trp Ala Asn Ile Leu Arg 85 90 Val Tyr Leu Thr Leu Lys Tyr Arg Phe Ser 105 106

<210> 2001 <211> 88 <212> PRT <213> Homo sapiens

<210> 2002 <211> 85 <212> PRT <213> Homo sapiens

<400> 2002 Met Arg Lys Leu Ile Ala Gly Leu Ile Phe Leu Lys Ile Trp Thr Cys 10 Thr Val Arg Thr Ser Thr Asp Leu Pro Gln Thr Glu Asp Cys Ser Gln 20 25 Cys Ile His Gln Val Thr Glu Ile Gly Gln Lys Val Ala Thr Val Leu 40 45 Leu Phe Tyr Gly Tyr Tyr Lys Tyr Thr Gly Thr Leu Lys Arg Thr Cys 55 60 Leu Tyr Asn Val Ile Leu Tyr Lys Val Tyr Ser Pro Gly Asn Asp Gln 70 Pro Asp Val Leu * 84

<210> 2003 <211> 46 <212> PRT <213> Homo sapiens

<210> 2004 <211> 51 <212> PRT <213> Homo sapiens

<400> 2004 Met Trp Leu Phe Ile Ala Ser Lys Cys Ile Phe Leu Leu Ile Val Pro Asn Phe Ile Phe Val Phe Trp Arg Lys Val Phe Ser His Asp Arg Leu 20 25 30 Asn Ile Ala Tyr Ser Phe Glu Leu Ser Ser Lys Tyr Ile Phe Ile Leu 35 Phe Ile 50 <210> 2005 <211> 66 <212> PRT <213> Homo sapiens <400> 2005 Met Val Glu Val Val Ser Leu Leu His Leu Tyr Ala Val Ala Cys Ala 1 5 10 Arg Lys Gly Pro Phe Pro Asn Thr Lys Asp Leu Ser Gly Trp Thr Pro 20 25 Ser Ser Gly Arg Glu Glu Leu Trp Lys Gly Lys Arg Ala Ala Ala Ala 40 45 Thr Arg Asn Pro Leu Val Leu Thr Gly Leu Gly Ser Pro Ser Ala Arg Leu * 65

<210> 2006 <211> 46 <212> PRT

<213> Homo sapiens

<210> 2007 <211> 87 <212> PRT <213> Homo sapiens

 $<\!400\!>$ 2007 . Met Pro Thr Leu Ala Lys Trp Ile Leu Ser Leu Ser Met Thr Ser Thr 1 5 10 15

<210> 2008

<211> 58

<212> PRT

<213> Homo sapiens

<400> 2008

<210> 2009

<211> 46

<212> PRT

<213> Homo sapiens

<400> 2009

 Met
 Leu
 Met
 Tyr
 Val
 Leu
 Pro
 Phe
 Cys
 Gly
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<210> 2010

<211> 235

<212> PRT

<213> Homo sapiens

<400> 2010

 Met Glu Leu Gly Cys Trp Thr Gln Leu Gly Leu Thr Phe Leu Gln Leu

 1
 5
 10
 15

 Leu Leu Ile Ser Ser Leu Pro Arg Glu Tyr Thr Val Ile Asn Glu Ala
 20
 25
 30

 Cys Pro Gly Ala Glu Trp Asn Ile Met Cys Arg Glu Cys Cys Glu Tyr

40 Asp Gln Ile Glu Cys Val Cys Pro Gly Lys Arg Glu Val Val Gly Tyr 55 Thr Ile Pro Cys Cys Arg Asn Glu Glu Asn Glu Cys Asp Ser Cys Leu Ile His Pro Gly Cys Thr Ile Phe Glu Asn Cys Lys Ser Cys Arg Asn 90 85 Gly Ser Trp Gly Gly Thr Leu Asp Asp Phe Tyr Val Lys Gly Phe Tyr 105 Cys Ala Glu Cys Arg Ala Gly Trp Tyr Gly Gly Asp Cys Met Arg Cys 120 Gly Gln Val Leu Arg Ala Pro Lys Gly Gln Ile Leu Leu Glu Ser Tyr 135 140 Pro Leu Asn Ala His Cys Glu Trp Thr Ile His Ala Lys Pro Gly Phe 150 155 Val Ile Gln Leu Arg Phe Val Met Leu Ser Leu Glu Phe Asp Tyr Met 170 165 Cys Gln Tyr Asp Tyr Val Glu Gly Cys Asp Gly Asp Asn Arg Asp Gly 185 His Ile Ile Lys Arg Val Cys Gly Asn Glu Arg Ala Ala Pro Ile His 200 205 Asn Ile Arg Ile Leu Thr Ser Arg Pro Phe Pro Leu Pro Gly Leu Ser 210 215 Lys Ile Leu Thr Gly Phe His Ala Pro Phe * 230

<210> 2011 <211> 61 <212> PRT <213> Homo sapiens

<210> 2012 <211> 107 <212> PRT <213> Homo sapiens

<210> 2013 <211> 67 <212> PRT <213> Homo sapiens

<210> 2014 <211> 59 <212> PRT <213> Homo sapiens

<210> 2015 <211> 55 <212> PRT <213> Homo sapiens

<400> 2015
Met Val Arg Leu Gln Val Leu Val Leu Val Phe Arg Val Val Gly Ser
1 5 10 . 15
Gln Gln Met Leu Arg Gln Gly Ala Ala Gly Ala Arg Ser His Arg Val

Leu Ala Ser Leu His Phe Gln His Gly Phe Gly Thr Phe His Thr Pro
35 40 45

Ala Arg Ala Gly Gly Ser Glu
50 55

<210> 2016 <211> 64 <212> PRT <213> Homo sapiens

<210> 2017 <211> 58 <212> PRT <213> Homo sapiens

<210> 2018 <211> 66 <212> PRT <213> Homo sapiens

Ile * 65

PATENT COOPERATION TREATY

PCT

DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

(PCT Article 17(2)(a), Rule 13ter.1(c) and 39)

Applicant's or agent's file reference		Date of mailing (day/month/year)
	IMPORTANT DECLARATION	0 m 11 15 2003
21272-018		8 7 JUN 2001
International application No.	International filing date (day/month/year)	(Earliest) Priority date (day/month/year)
increasional application 140.	international filing take (tary/morally year)	(Lancst) i nonty trace (ausymorani year)
PCT/US01/02687	25 January 2001 (25.01.2001)	25 January 2000 (25.01.2000)
International Patent Classification (IPC) or both national classification and IPC		
IPC(7): C12P 21/06 and US Cl.: 435/69.1		
Applicant		
HYSEQ, INC.		
This International Searching Authority hereby declares, according to Article 17(2)(a), that no International search report		
will be established on the international application for the reasons indicated below.		
1. The subject matter of the international application relates to:		
b mathematical theories		
c. plant varieties.		
d. animal varieties.		
e. essential biological processes for the production of plants and animals, other than microbiological processes		
and the products of such processes.		
f schemes, rules or methods of doing business.		
g. schemes, rules or methods of performing purely mental acts.		
h. schemes, rules or methods of playing games.		
i. methods for treatment of the human body by surgery or therapy.		
j. methods for treatment of the animal body by surgery or therapy.		
k. diagnostic methods practised on the human or animal body.		
1 mere presentations of information.		
m computer programs for which this International Searching Authority is not equipped to search prior art.		
2. The failure of the following parts of the international application to comply with prescribed requirements prevents a		
meaningful search from bein	· —	7
the description	the claims	the drawings
. [7]		to the second amounted for its Annow C
3. The failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions prevents a meaningful search from being carried out:		
(
the written form has not been furnished or does not comply with the standard.		
the computer readable form has not been furnished or does not comply with the standard.		
4. Further comments:		
4. Further comments:		
		•
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Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Authorized officer Authorized officer		
Box PCT Young J. Kim		
Washington, D.C. 20231		

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